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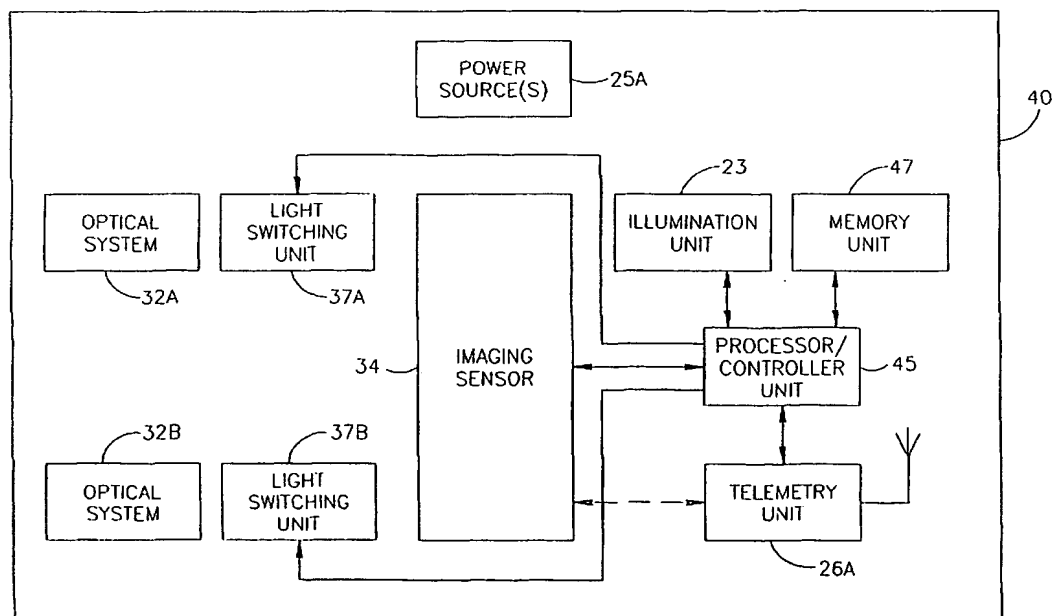
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(54) Title: APPARATUS AND METHODS FOR IN VIVO IMAGING



(57) Abstract: An in vivo imaging device and method, the device including at least one illumination source; at least one image sensor; and at least two optical systems. The optical systems have different depths of focus. A first and second image are focused onto the image sensor.

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APPARATUS AND METHOD FOR IN VIVO IMAGING

FIELD OF THE INVENTION

The present invention relates generally to in vivo imaging devices, and more specifically to in vivo imaging devices having extended depth of field or
5 variable depth of field.

BACKGROUND OF THE INVENTION

Devices and methods for performing in-vivo imaging of passages or cavities within a body are known in the art. Such devices may include, inter alia, various endoscopic imaging systems and devices for performing imaging in
10 various internal body cavities.

Reference is now made to Fig. 1 which is a schematic diagram illustrating an embodiment of an autonomous in-vivo imaging device. The device 10A typically includes an optical window 21 and an imaging system for obtaining images from inside a body cavity or lumen, such as the GI tract. The imaging
15 system includes an illumination unit 23. The illumination unit 23 may include one or more light sources 23A. The one or more light sources 23A may include a white light emitting diode (LED), or any other suitable light source, known in the art. The device 10A includes a CMOS imaging sensor 24, which acquires the images and an optical system 22 which focuses the images onto the CMOS
20 imaging sensor 24.

The optical system 22 may include one or more optical elements (not shown), such as one or more lenses (not shown), one or more composite lens assemblies (not shown), one or more suitable optical filters (not shown), or any other suitable optical elements (not shown) adapted for focusing an image of
25 the GI tract on the imaging sensor as is known in the art and disclosed hereinabove with respect to the optical unit 22 of Fig. 1. The optical system 22 may be attached to, or mounted on, or fabricated on or disposed adjacent to the imager light sensitive pixels (not shown) as is known in the art.

The illumination unit 23 illuminates the inner portions of the body lumen or body cavity (such as, for example the gastrointestinal cavity) through an optical window 21. Device 10A further includes a transmitter 26 and an antenna 27 for transmitting image signals from the CMOS imaging sensor 24, and one or more
5 power sources 25. The power source(s) 25 may be any suitable power sources such as but not limited to silver oxide batteries, lithium batteries, or other electrochemical cells having a high energy density, or the like. The power source(s) 25 may provide power to the electrical elements of the device 10A.

Typically, in the gastrointestinal application, as the device 10A is
10 transported through the gastrointestinal (GI) tract, the imager, such as but not limited to the multi-pixel CMOS sensor 24 of the device 10A acquires images (frames) which are processed and transmitted to an external receiver/recorder (not shown) worn by the patient for recording and storage. The recorded data may then be downloaded from the receiver/recorder to a computer or
15 workstation (not shown) for display and analysis. During the movement of the device 10A through the GI tract, the imager may acquire frames at a fixed or at a variable frame acquisition rate. For example, the imager (such as, but not limited to the CMOS sensor 24 of Fig. 1) may acquire images at a fixed rate of two frames per second (2 Hz). Other different frame rates may also be used,
20 depending, inter alia, on the type and characteristics of the specific imager or camera or sensor array implementation that is used, and on the available transmission bandwidth of the transmitter 26. The downloaded images may be displayed by the workstation by replaying them at a desired frame rate. This way, the expert or physician examining the data is provided with a movie-like
25 video playback, which may enable the physician to review the passage of the device through the GI tract.

Typically, the device 10A or a similar autonomous in vivo imaging device is propelled through the GI tract by the natural action of peristalsis. When the device 10A or a similar autonomous in vivo imaging device is used for imaging,
30 some of the acquired images may be out of focus. Additionally in some of the acquired images, a part or parts of the acquired image may be out of focus because of a possibly limited depth of field obtainable by the optical system 22.

For example, if the optical system 22 includes a lens (not shown in detail) having a limited depth of field, and the imaged object (such as, for example, the wall of the GI tract) or a portion thereof was not disposed at a distance which is within the depth of field range of the lens, the acquired image or parts thereof may be blurred or not depicted sharply.

It may therefore be desirable to decrease the number or the percentage of acquired images which are not acceptably focused or which have a reduced sharpness or detail due to out of focused imaging.

SUMMARY

Embodiments of the present invention provide an in vivo imaging device and method. In one embodiment, a device includes an illumination source; an image sensor; and at least two optical systems. The optical systems have different depths of focus. A first and second image are focused onto the image sensor.

BRIEF DESCRIPTION OF THE DRAWINGS

The invention is herein described, by way of example only, with reference to the accompanying drawings, in which like components are designated by like reference numerals, wherein:

Fig. 1 is a schematic diagram illustrating an embodiment of an autonomous in-vivo imaging device;

Fig. 2 is a schematic cross-sectional view of part of an in-vivo imaging device having two optical systems and a single imager, in accordance with an embodiment of the present invention;

Fig. 2A is a schematic front view of the surface of the imaging sensor of the device illustrated in Fig. 2, schematically illustrating non-overlapping images projected on the surface of the imaging sensor;

Fig. 3 is a schematic cross-sectional view of part of an in-vivo imaging device having two shuttered optical systems and a single imager, in accordance with another preferred embodiment of the present invention;

5 Fig. 3A is a schematic front view of the surface of the imaging sensor of the device illustrated in Fig. 3, schematically illustrating the degree of overlap of images which may be projected on the surface of the imaging sensor if both shutters are opened; and

Fig. 4 is a schematic functional block diagram useful in understanding the components of the device partially illustrated in Fig. 3, according to an
10 embodiment of the invention.

DETAILED DESCRIPTION OF THE INVENTION

One approach which may be used to solve the image sharpness problem of in vivo imaging devices is to use an optical system such as, but not limited to,
15 a lens or a lens assembly having a wide depth of field range. Typically, wide angle lenses may be used. Such lenses may be compound multi-element lenses or other lens types.

Another approach may be to use a plurality of lenses within the same imaging device.

20 Reference is now made to Fig. 2 which is a schematic cross-sectional view of part of an in-vivo imaging device having two optical systems and a single imager, in accordance with an embodiment of the present invention.

The in vivo imaging device 30 (only part of which is illustrated in Fig. 2) includes an imaging sensor 34. The imaging sensor 34 may be a CMOS pixel
25 array sensor similar to the CMOS imaging sensor 24 of Fig. 1, or may be any other type of suitable imaging sensor known in the art; for example, a CCD may be used.

The device 30 includes a housing 31 and an optical window 21A attached to the housing 31. If the device 30 is implemented as a swallowable capsule,
30 the housing 31 may be, for example, a capsule-like housing, as disclosed in

detail in U.S. patent 5,604,531 to Iddan et al., and/or WO 01/65995 to Glukhovsky et al. The system and method of the present invention may be used with other swallowable devices. If the device 30 is implemented as, for example, an endoscope-like device or a catheter-like device, the housing 31
5 may be an extended elongated flexible device (the extended device is not shown in its entirety for the sake of clarity of illustration). Such devices may be shaped as, for example, elongated flexible devices for insertion into a body cavity or body lumen of a human patient or an animal.

The optical window 21A may be a transparent optical dome as disclosed
10 in WO 00/76391 to Glukhovsky et al. but may be any type of suitable optical window.

The device 30 may include an illumination unit 33 which may include one or more light sources 33A. The light sources 33A may be "white" LED sources as disclosed in WO 01/65995 to Glukhovsky et al. but may also be any other
15 light sources suitable for providing illumination for in vivo imaging, such as but not limited to the light sources disclosed in U.S. patent 5,604,531 to Iddan et al. The device 30 may include two optical systems 32A and 32B. The optical systems 32A and 32B may be attached to or mounted in a baffle 35, or otherwise suitably attached to the housing 31 of the device 30 or to any other
20 suitable structure included in the device 30 (such as for example, the imaging sensor 34). The baffle 35 may be shaped to prevent light emitted from the light sources 33A from directly reaching the optical systems 32A and 32B, while allowing light reflected from the imaged object, such as but not limited to, the intestinal wall (not shown) to reach the optical systems 32A and 32B. The
25 device 30 may be of other configurations and include other combinations of components. For example, the baffle 35 need not be used.

The imaging sensor 34 may be disposed adjacent to the optical systems 32A and 32B, or may be also attached thereto.

The optical systems 32A and 32B may be typically single lenses,
30 composite (multi-element) lenses, or any suitable combination of optical elements which are suitable for forming images of the imaged object, (such as

but not limited to, the intestinal wall) and projecting the images onto the surface of the imaging sensor 34.

The depth of field range of the optical system 32A is different than the depth of field range of the optical system 32B. For example, the focal length of the optical system 32A may be longer than the focal length of the optical system 32B. Each of the optical systems 32A and 32B projects an image on the surface of the imaging sensor 34. According to one embodiment, the images projected by the optical systems 32A and 32B do not overlap.

Thus, after the pixels of the imaging sensor 34 are scanned (read out) and the data is transmitted to an outside receiver/recorder as disclosed hereinabove, the resulting image may have two typically non-overlapping parts. One image part corresponds to the image projected on the imaging sensor by the optical system 32A, and the other image part corresponds to the image projected on the imaging sensor by the optical system 32B.

Because of the different focal length of the two different optical systems 32A and 32B, even if one image part is not properly focused the other image part may be in focus due to the larger depth of field range of the optical system having the shorter focal length. Thus, the chance of obtaining at least one acceptably focused image is increased relative to the chance in a similar in vivo imaging device which includes only a single optical system 32A (for example, having a larger focal length and a narrower depth of focus) or which may include a plurality of optical systems.

Reference is now made to Fig. 2A which is a schematic front view of the surface of the imaging sensor of the device illustrated in Fig. 2, schematically illustrating non-overlapping images projected on the surface of the imaging sensor.

In Fig. 2A, the surface 34A represents a top view of the surface of the entire imaging sensor 34. The surface 34B schematically represents the imager area part comprising the light sensitive pixels (the pixels are not shown in detail). The part of the imager surface surrounding the area 34B may include the support circuitry for performing readout of the pixels and other electronic support circuitry such as clocking circuitry and the like, as is known in the art.

The circular areas 36A and 36B schematically represent the area of the images projected on the surface 34B by the optical systems 32A and 32B, respectively. The areas 36A and 36B do not overlap.

5 It will be appreciated by those skilled in the art that while the device 30 of Fig. 2 may provide a solution to the depth of focus problem, it may not make the best use of the light sensitive pixel areas 34B available in the imaging sensor 34. This may be due to the requirement for non-overlapping of the projected image areas 36A and 36B.

10 Reference is now made to Fig. 3, which is a schematic cross-sectional view of part of an in-vivo imaging device having two optical systems and a single imager, in accordance with another preferred embodiment of the present invention.

The in vivo imaging device 40 includes an imaging sensor 34. The imaging sensor 34 may be a CMOS pixel array sensor similar to the CMOS
15 imaging sensor 24 of Fig. 1, or may be any other type of suitable imaging sensor known in the art.

The device 40 includes a housing 31 and an optical window 21A attached to the housing 31, as disclosed hereinabove and illustrated in Fig. 2. If the device 40 is implemented as a swallowable capsule, the housing 31 may be a
20 capsule-like housing, as disclosed in detail in U.S. patent 5,604,531 to Iddan et al., and/or in WO 01/65995 to Glukhovsky et al.. If the device 40 is implemented as an endoscope-like device or a catheter-like device, the housing 31 may be an extended elongated flexible device (the extended device is not shown in it's entirety for the sake of clarity of illustration). Such devices may be
25 shaped as elongated flexible devices for insertion into a body cavity or body lumen of a human patient or an animal.

The optical window 21A may be a transparent optical dome as disclosed in WO 00/76391 to Glukhovsky et al. but may be any type of suitable optical window.

30 The device 40 may include an illumination unit 33, which may include one or more light sources 33A, as disclosed hereinabove. The light sources 33A may be "white" LED sources but may also be any other light sources suitable for

providing illumination for in vivo imaging, such as but not limited to the light sources disclosed in U.S. patent 5,604,531 to Iddan et al., and in The device 40 may include two optical systems 32A and 32B. The optical systems 32A and 32B may be attached to or mounted in a baffle 35A, or otherwise suitably
5 attached to the housing 31 of the device 40 or to any other suitable structure included in the device 40 (such as, for example, to the imaging sensor 34). The baffle 35A may be shaped to prevent light emitted from the light sources 33A from directly reaching the optical systems 32A and 32B, while allowing light reflected from the imaged object, such as but not limited to, the intestinal wall
10 (not shown) to reach the optical systems 32A and 32B.

The imaging sensor 34 may be disposed adjacent to the optical systems 32A and 32B, or may be also attached thereto.

The optical systems 32A and 32B may be single lenses, composite (multi-element) lenses, or any suitable combination of optical elements which
15 are suitable for forming images of the imaged object, (such as but not limited to, the intestinal wall) and projecting the images onto the surface of the imaging sensor 34.

The depth of field range of the optical system 32A is different than the depth of field range of the optical system 32B. For example, the focal length of
20 the optical system 32A may be longer than the focal length of the optical system 32B. Each of the optical systems 32A and 32B projects an image on the surface of the imaging sensor 34. In contrast to the arrangement of the optical components illustrated in Fig. 2, the images projected by the optical system 32A and 32B on the imaging sensor 34 of the device 40 may overlap.

25 The device 40 further includes two controllable light switching units 37A and 37B. According to an embodiment of the invention the controllable light switching unit 37A is interposed between the optical system 32A and the imaging sensor 34. The controllable light switching unit 37B is interposed between the optical system 32B and the imaging sensor 34. Preferably, but not
30 necessarily, the controllable light switching units 37A and 37B may be electro-optical devices which may be electrically (or magnetically) controlled to block or enable the passage of light therethrough. For example, the controllable

light switching units 37A and 37B may be electrically controllable liquid crystal shutters (LCS), electrically controllable ferroelectric optical shutters, high-speed electrolytic optical shutters, electro-optical shutters based on the Kerr and Pockels effects, or optical shutter devices based on ferroelectric films or ferroelectric liquids or on ferroelectric crystals or any other suitable electro-optical or magneto-optical shutter known in the art. The controllable light switching units 37A and 37B may also be any suitable controllable electro-mechanical shutter devices known in the art.

In operation, the controllable light switching units 37A and 37B may be used to provide exposure of the imaging sensor 34 to light projected by a selected one of the optical systems 32A and 32B. This selection is typically needed due to the at least partial overlap of the images projected on the surface of the imaging sensor 34 as shown hereinafter.

Reference is now briefly made to Fig. 3A which is a schematic front view of the surface of the imaging sensor of the device illustrated in Fig. 3, schematically illustrating the degree of overlap of images which may be projected on the surface of the imaging sensor if both light switching units (shutters) are opened, according to an embodiment of the invention.

In Fig. 3A, the surface 34A represents a top view of the surface of the entire imaging sensor 34. The surface 34B schematically represents the imager area part comprising the light sensitive pixels (the pixels are not shown in detail). The part of the imager surface surrounding the area 34B may include the support circuitry for performing readout of the pixels and other electronic support circuitry such as clocking circuitry and the like, as is known in the art. The partially overlapping circular areas 39A and 39B schematically represent the area of the images projected on the surface 34B by the optical systems 32A and 32B, respectively. The areas 39A and 39B overlap. The hatched area 39C represents the area of overlap between the image projected on the surface 34B by the optical system 32A and the image projected by the optical system 32B on the surface 34B.

It is noted that the image overlap shown may occur only in a situation in which both of the light switching units 37A and 37B are switched to allow

passage of the light coming from the corresponding optical systems 32A and 32B to reach the surface 34B of the imaging sensor 34. This typically does not normally occur during operation of the device 40, since the purpose of the light switching units 37A and 37B is typically to prevent the simultaneous exposure of the surface 34B to light projected from both optical systems 32A and 32B.

Thus, in operation, during active imaging, the illuminating unit 23 may be switched on to provide illumination for an imaging cycle. An imaging cycle may comprise a first imaging time period in which the light switching unit 37A may be controllably switched on to allow the light collected by the optical system 32A to be projected on the surface 34B of the imaging sensor 34 while during the same first imaging time period the light switching unit 37B is switched off such that light projected from the optical system 32B is blocked by the light switching unit 37B and does not reach the surface 34B of the imaging sensor 34. In the first time period a first image projected by the optical system 32A is acquired.

After the first image projected by the optical system 32A is acquired by scanning (readout) of the pixels of the imager 34, the image data may be stored in a memory device (not shown) included in the device 40 for later transmission, or may be directly transmitted to an external receiver/recorder as disclosed hereinabove. After the acquired image data is stored in a memory device or is transmitted to an external receiver/recorder, a second imaging time period may be started. During the second imaging time period, the light switching unit 37B may be controllably switched on to allow the light collected by the optical system 32B to be projected on the surface 34B of the imaging sensor 34 while during the same second time period the light switching unit 37A is switched off such that light projected from the optical system 32A is blocked by the light switching unit 37B and does not reach the surface 34B of the imaging sensor 34. In the second imaging time period a second image projected by the optical system 32B is acquired. After the end of the imaging cycle, which may be the time of termination of the second imaging time period, the illumination unit 23 may be turned off.

After the second image projected by the optical system 32B is acquired by scanning (readout) of the pixels of the imager 34, the image data may be

stored in a memory device (not shown) included in the device 40 for later transmission, or may be directly transmitted to an external receiver/recorder as disclosed hereinabove.

5 If the first image data and the second image data have been stored in a memory device (not shown) the data of the first and the second acquired images may be transmitted to the receiver/recorder. Alternatively, the first and the second acquired images may be transmitted after the second imaging time period is terminated.

10 The data transmitted may be stored in the receiver/recorder device (not shown) for later processing and display. Thus, each imaging cycle of the device 40 may yield two different images of approximately (but not necessarily exactly) the same imaged object.

15 It will be appreciated that while the imaging method used in the device 30 (Fig. 2) may yield two images of acquired simultaneously within a single imaging cycle through two different optical systems having a different depth of focus range, the imaging method used in the device 40 (Fig. 3) may yield two images sequentially acquired through two different optical systems having a different depth of focus range. If the device 40 does not move (remains stationary) within the duration of the imaging cycle, the images may represent approximately the same object or imaged area (which may however be imaged with different field of view due to the different field of view of the optical systems 32A and 32B). If the imaging device 40 moves during the imaging cycle, the first and second images acquired may not show the same object or the object may be shifted in the second image relative to the first image. The degree of the shift may depend, inter alia, on the duration of the imaging cycle, the velocity of the imaging device 40 relative to the imaged object (such as, for example, the intestinal wall), the distance to the imaged object, and the focal length of the optical systems 32A and 32B.

25 Preferably, the duration of the imaging cycle should be short to prevent or reduce image blurring or smearing (which may occur due to device movement in both of the devices 30 and 40), and to reduce or prevent shift of the imaged

objects which may occur in the device 40 due to the sequential acquisition of the first image and the second image in an imaging cycle.

Thus, while the device 30 has the advantage of not being subject to the image shift as disclosed hereinabove, it may have lower image resolution since two images is simultaneously acquired on the surface 34B of the imaging sensor 34 with no image overlap which reduces the number of pixels within each of the two images. The device 40 may have the advantage of higher resolution for each image because only one image may be acquired at the same time, allowing better use of the available surface 34B of the imaging sensor 34 in that each of the two images which are acquired sequentially may have a higher pixel number than the images acquired simultaneously in the device 30. The device 40 may however be more susceptible to image shifting caused by movement of the device 40 within the duration of a single imaging cycle. This image shift may be reduced by reducing the duration of the imaging cycle or by reducing the time of acquisition of each of the two images within the imaging which may be achieved, inter alia, by using an imaging sensor having a high sensitivity.

Because of the different focal length of the two different optical systems 32A and 32B, even if one image part is not properly focused the other image part may be in better focus due to the larger depth of field range of the optical system having the shorter focal length. Thus, the chance of obtaining at least one acceptably focused image within a single imaging cycle of in-vivo imaging devices 30 and 40 is increased relative to the chance in a similar in vivo imaging device which includes only a single optical system.

It is noted that while the overlap (illustrated in Fig. 3A) between the areas 39A and 39B of Fig. 3A is partial, the optical systems 32A and 32B of the device 40 may be configured such that the areas 39A and 39B may fully overlap. This may be achieved if desired by suitable configuring of the optical systems 32A and 32B of the device 40 as is known in the art or by introducing additional optical elements (not shown) to ensure overlap of the two areas 39A and 39B. The advantage of full overlap is that such a design enables a better utilization of

the surface 34B which includes the light sensitive pixel elements, and may achieve a higher resolution for both images without reducing the image size.

Reference is now made to Fig. 4, which is a schematic functional block diagram useful in understanding the operation of the device illustrated in Fig. 3 and, possibly with some modifications (e.g., possibly removal of units 37a and 37b) Fig. 2.

The device 40 may include one or more power sources 25A for supplying power to the components included in the device 40. The device 40 may include a processor/controller unit 45. The processor/controller unit 45 may be suitably connected to an imaging sensor 34 for controlling the operation of the imaging sensor 34 and for (optionally) receiving the image data from the imaging sensor 34. The imaging sensor 34 may be (optionally) suitably connected to a telemetry unit 26A. The telemetry unit 26a may receive the data or signals read out from the imaging sensor 34 and transmit the signals or data (with or without further processing) to a receiver/recorder (not shown in Fig. 4) as is disclosed in detail hereinabove. The telemetry unit 26A may operate similar to the transmitter 26 coupled to the antenna 27 of Fig. 1. The device 40 may further include two different optical systems 32A and 32B having different depth of focus range as disclosed hereinabove and illustrated in Fig. 3.

A light switching unit 37A may be interposed between the optical system 32A and the light sensitive surface of the imaging sensor 34. Another light switching unit 37B may be interposed between the optical system 32B and the light sensitive surface of the imaging sensor 34. The light switching units 37A and 37B are suitably connected to the controller/processor unit 45 and may receive control signals therefrom for switching the light switching units 37A and/or 37B on and off. The switching signals may be digital signals, or may be analog voltage signals produced by a suitable interface unit (not shown), but may however be any other suitable control signals known in the art for switching the switching units on or off. In alternate embodiments, control may be provided in other manners, using other components or combinations of components. For example, the telemetry unit 26A may provide control.

The device 40 includes an illuminating unit 23 as disclosed in detail hereinabove and illustrated in Figs. 1, 2, and 3. The illumination unit may be suitably connected to the processor controller unit 45 for receiving control signals therefrom. The processor controller unit 45 may control the operation of the illumination unit 23 by switching it on or off as necessary.

The device 40 may (optionally) include a memory unit 47 suitably connected to the processor/controller unit. The memory unit 47 may be (optionally) used to store acquired image data which is read out from the imaging sensor 34.

It is noted that the imaging sensor 34 may or may not include further processing circuitry (not shown) which may be used for performing imaging various support functions such as synchronization and clocking functions, pixel scanning functions and other functions associated with imaging sensor which are known in the art of imaging sensors. Additionally, the imaging sensor 34 may also include analog to digital converting circuitry (not shown in detail) for providing digital signals to the processor/controller unit 45. Alternatively, a separate analog to digital converter (not shown) may (optionally) couple the imaging sensor 34 to the processor/controller unit 45.

For performing an imaging cycle, the device 40 may be operated as follows. At the beginning of each imaging cycle, the processor/controller unit 45 may switch on the light switching unit 37A and may simultaneously switch off the light switching unit 37B to allow light focused by the optical system 32A to be projected onto the surface of the imaging unit 34 while blocking the light focused by the optical system 32B from reaching the surface of the imaging unit 34. Simultaneously with the switching of the light switching unit 37A on, the processor/controller unit 45 may switch on the illumination unit 23 to illuminate the imaged object (not shown). After the end of the first imaging period, the processor/controller unit 45 may store the acquired first image in the memory unit 47 or may alternatively control the transmission of the acquired image to a receiver/recorder (not shown).

If the first image data is transmitted by the telemetry unit 26A after image acquisition, the illumination unit 23 may be turned off for the duration of

the transmission time period to conserve power and to reduce the current drain from the power source(s) 25A. In such a case, the processor/controller unit 45 may switch the illumination unit on again after the data has been transmitted. Simultaneous with switching the illumination unit on the processor/controller unit 5 45 may also switch on the light switching unit 37B and may switch off the light switching unit 37A. In this way the light focused by the optical system 32B is now allowed to reach the surface of the imaging sensor 34 for the duration of a second imaging time period and a second image is acquired. After the termination of the second imaging period, the illumination unit 23 may be turned 10 off by the processor/controller unit 45. The second acquired image may then be transmitted by the telemetry unit 26A as disclosed hereinabove. The device 40 may then be ready for another imaging cycle.

If the device 40 has image storage capacity (such as by including the memory unit 47 of Fig. 40), the first acquired image may be stored in the 15 memory unit 47 while the pixels readout of the imaging sensor 34 is performed. In such a case, the processor/controller unit 45 may turn off the illuminating unit 23 at the end of the first imaging period and may turn the illuminating unit 23 on again after the image data readout has been completed and the imaging sensor has been reset for enabling the acquiring of a second image. In such a case, 20 the data of the second image acquired within the imaging cycle may be also stored in the memory unit 47 and the data of the first acquired image and the second acquired image may be transmitted by the telemetry unit 26A. The stored data of the first and second acquired images may be transmitted sequentially. Other forms of transmission may however also be possible. For 25 example, it may be possible to transmit data from the first and second images in an interleaved manner.

It is noted, however, that if the device 40 includes a memory device such as the memory unit 47, it may also be possible to start transmitting the stored data of the first acquired image before the acquisition of the second image has 30 been completed. For example, the telemetry unit 26A may be controlled to start the transmission of the first image data as soon as some data for the first image is stored in the memory device 47. Such transmission may be controlled and

timed by the processor/controller 45. The advantage of this transmission method is that it may enable transmission of the first image data while acquisition of the second image data is being performed, which may enable the repeating of the imaging cycles at a higher frequency than the frequency possible in the method in which both the first and the second image data are transmitted sequentially only after the acquisition of both the first and the second images has been completed.

It is noted that the memory unit 47 may be, a random access memory unit (RAM) but any other suitable type of memory device or unit known in the art may be used. The memory unit 47 may be a separate memory unit suitably coupled or connected to the processor/controller unit 45 or may be a memory unit included in the same integrated circuit (IC) on which the processor/controller unit 45 is made, which may obviate the need for connecting a separate memory unit to the processor/controller unit 45.

It is further noted that it may be possible to fabricate some or all of the electronic circuitry or electronic components illustrated in Fig. 4 on the same Integrated circuit. Thus, for example, it may be possible to fabricate two or more of the processor/controller 45, the memory unit 47, and the telemetry unit 26A on the same IC to save space and to simplify assembly of the device 40. It may also be possible to fabricate one or more of the processor/controller 45, the memory unit 47, and the telemetry unit 26A on the same IC on which the imaging sensor 34 is fabricated.

It will be appreciated by those skilled in the art, that while the devices 30 and 40 are devices that include two optical systems 32A and 32B, other embodiments of the invention may be constructed, which include more than two optical systems (not shown). In accordance with one embodiment of the invention, such devices may have no light switching units and may operate by simultaneous acquisition of more than two non-overlapping images. In accordance with another embodiment of the invention, such devices may have light switching units and may operate by sequential imaging of potentially overlapping images.

It may also be possible to use a combination of the operating methods disclosed hereinabove in an imaging device having a plurality of optical systems. For example, in accordance with an embodiment of the present invention the in vivo imaging device may include three optical systems (not shown). A first optical system and a second optical system of the three optical systems may project potentially overlapping or partially overlapping images on the surface of the imaging sensor (not shown). These first and second optical systems may be associated with two light switching elements (not shown) which may be used as disclosed hereinabove to sequentially acquire the two images sequentially projected on the imaging sensor. The third optical system (not shown) may project a third image on the surface of the imaging unit. This third image may be non-overlapping with the first and the second images. The third image may thus be acquired simultaneously with the first image projected by the first optical system, or (alternatively) simultaneously with the second image projected by the second optical system.

While the invention has been described with respect to a limited number of embodiments, it will be appreciated that many variations, modifications and other applications of the invention may be made which are within the scope and spirit of the invention.

CLAIMS

1. An in vivo imaging device comprising
at least one illumination source;
5 at least one image sensor; and
at least two optical systems, said optical systems having different depths
of focus and said optical systems capable of focusing at least a
first and second image onto said at least one image sensor.
2. The device according to claim 1 wherein the at least first and at least
10 second image do not overlap.
3. The device according to claim 1 wherein the at least first and at least
second image at least partially overlap.
4. The device according to claim 1 further comprising at least two light
switching units.
- 15 5. The device according to claim 3 further comprising at least two light
switching units.
6. The device according to claim 4 or 5 wherein at least a first light switching
unit is switched on during a first imaging time period and wherein at least a
second light switching unit is switched on during a second imaging time period.
- 20 7. A device according to claim 6 wherein the first light switching unit is
switched off after the first imaging time period and wherein the second light
switching unit is switched off after the second imaging time period.
8. The device according to claim 4 further comprising a controller for
controlling the switching of at least one light switching unit.
- 25 9. The device according to claim 1 further comprising a memory device.
10. The device according to claim 1 further comprising a transmitter.

11. The device according to claim 1, wherein the device is a swallowable capsule.
12. An in vivo imaging device comprising
at least one illumination source;
5 at least one image sensor having a light sensitive surface; and
at least two optical systems, said optical systems having different depths
of focus and said optical systems capable of focusing at least a
first and second image onto said the light sensitive surface.
13. The device according to claim 12 further comprising at least two light
10 switching units.
14. The device according to claim 13 wherein at least one light switching unit
is interposed between at least a first optical system and the light sensitive
surface.
15. The device according to claim 12, wherein the device is a swallowable
15 capsule.
16. A method for obtaining in vivo images, the method comprising the steps
of:
focusing light remitted from an in vivo site onto at least one image
sensor through at least a first optical system; and
20 focusing light remitted from the in vivo site onto at least one image
sensor through at least a second optical system.
17. The method according to claim 16 further comprising the step of reading
out of image data into a memory device.
18. The method according to claim 16 further comprising transmitting image
25 data from the memory device to an external receiving unit.
19. The method according to claim 16 further comprising a step of
transmitting image data to an external receiving unit.

20. A method for obtaining in vivo images, the method comprising the steps of:

5 focusing light remitted from an in vivo site onto at least one image sensor through at least a first optical system during a first imaging time period; and

focusing light remitted from the in vivo site onto at least one image sensor through at least a second optical system during a second imaging time period.

10 21. The method according to claim 20 further comprising the step of reading out of image data into a memory device.

22. The method according to claim 20 further comprising transmitting image data from the memory device to an external receiving unit.

23. The method according to claim 20 further comprising a step of transmitting image data to an external receiving unit.

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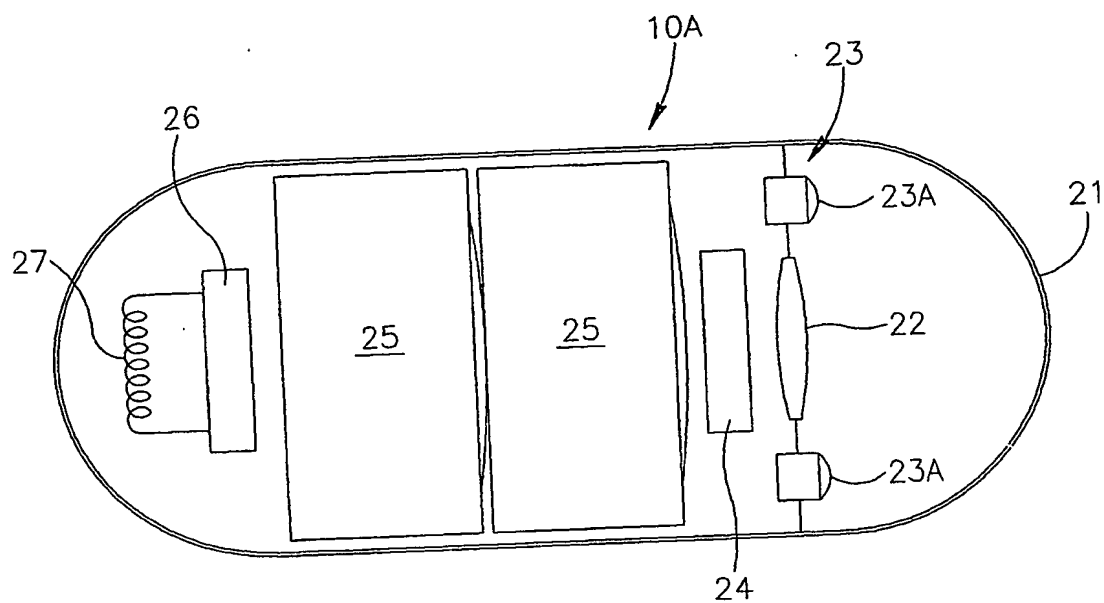


FIG.1

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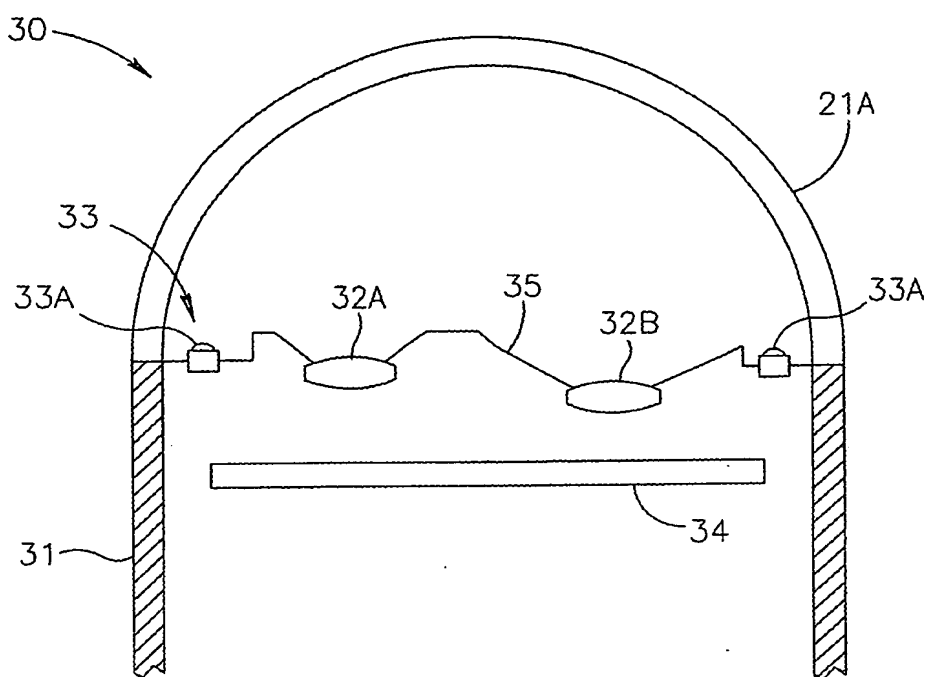


FIG. 2

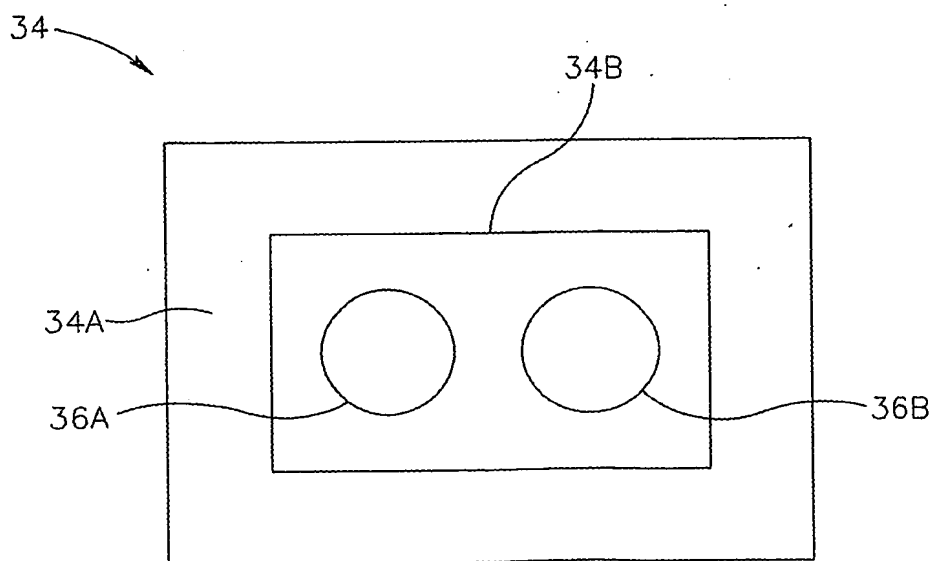


FIG. 2A

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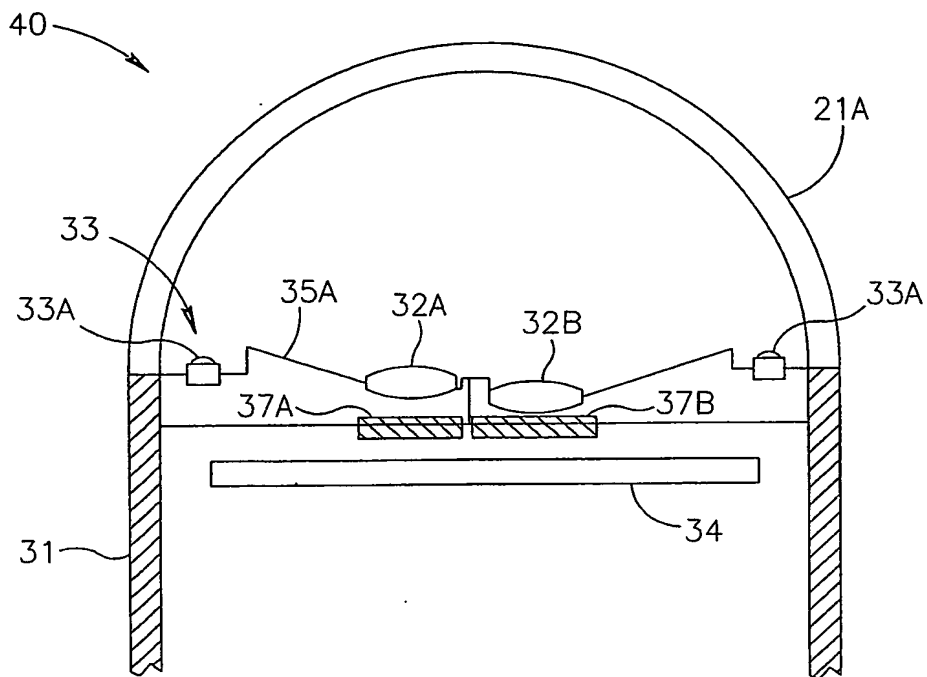


FIG. 3

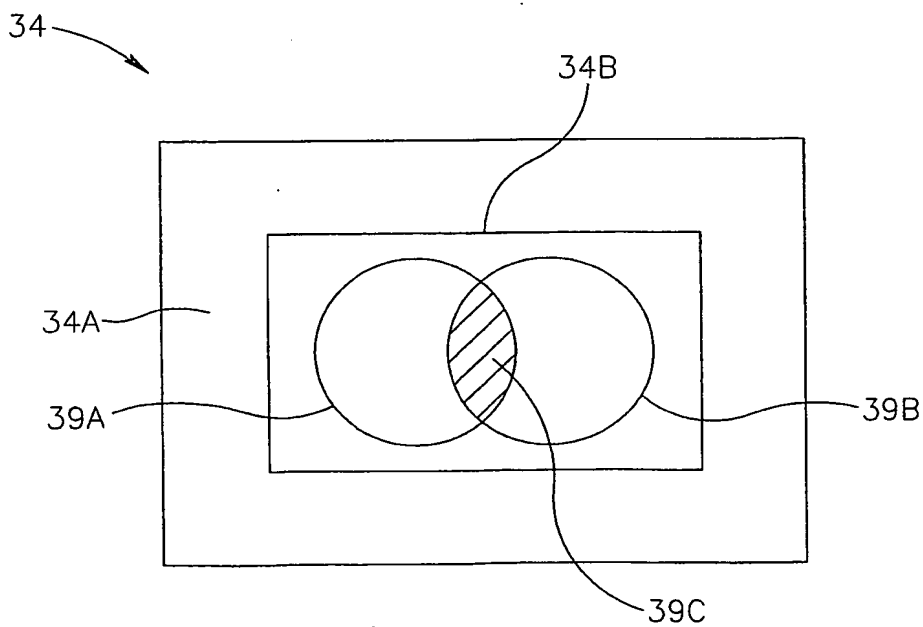


FIG. 3A

4/4

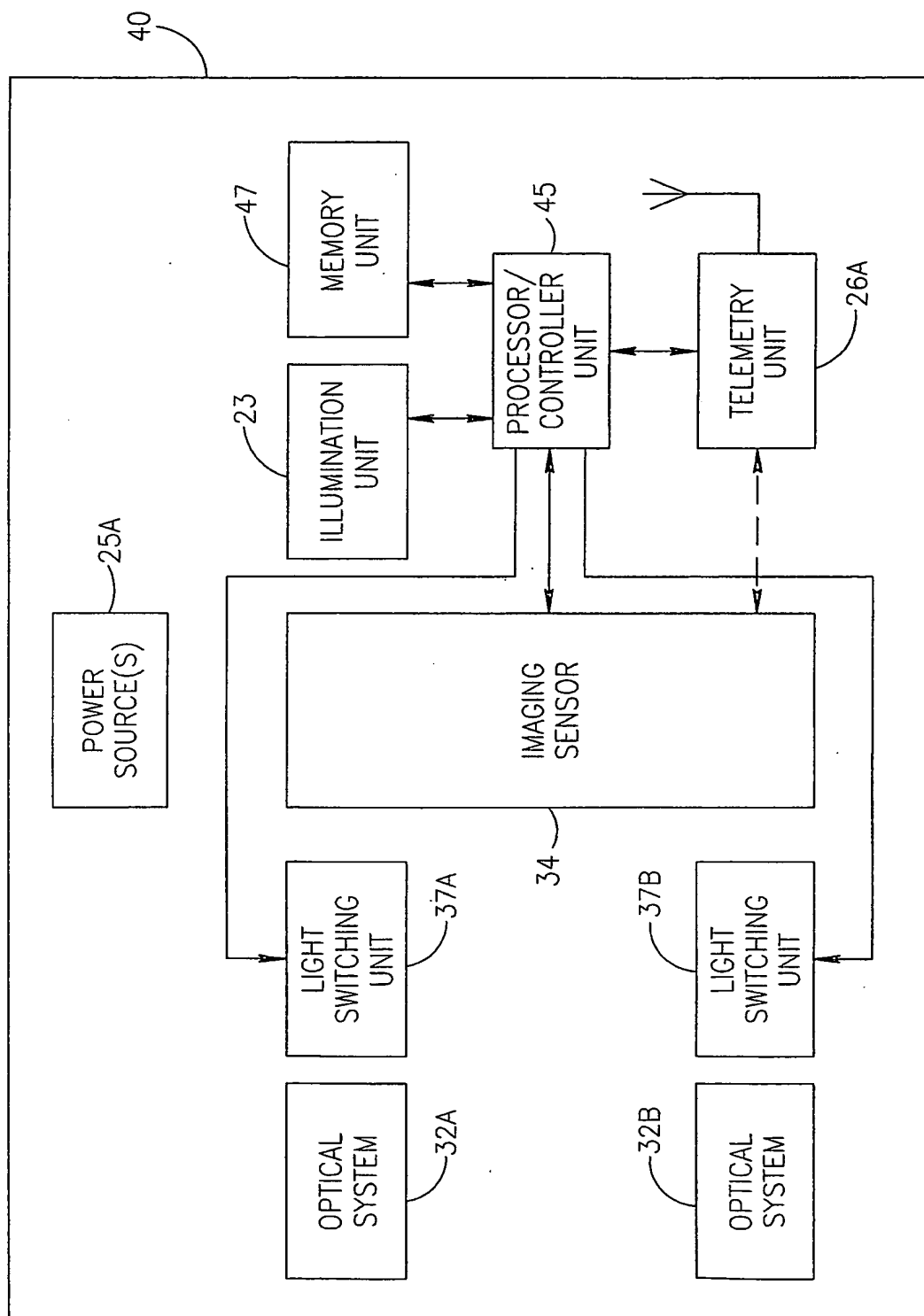


FIG. 4

DEVICE, SYSTEM AND METHOD FOR SELECTIVE ACTIVATION OF IN VIVO SENSORS

5

FIELD OF THE INVENTION

The present invention relates to the field of in vivo devices. More specifically, the present invention relates to a device, system and method for selectively activating or
10 altering the operational mode of an in vivo device, for example, in response to in vivo conditions.

BACKGROUND OF THE INVENTION

15

Certain in vivo devices may be introduced into a body in a location remote to the area where their sensing, diagnosing or other functions may be performed. For example, an in vivo device for imaging areas of the small intestine may be introduced into a body through the mouth and pass through the stomach and other parts of the gastrointestinal
20 (GI) tract by way of peristalsis until reaching the small intestine. Similarly, an in vivo device may be introduced into a body wherein the location of an area of interest or of a suspected pathology may be unknown or uncertain, thereby necessitating that an in vivo device pass from its point of introduction and locate the area of pathology where its sensing functions or other functions may be required for diagnosing pathologies or
25 performing other functions.

In vivo devices such as sensors are generally configured to capture sensory data on a fixed schedule that may be set or programmed into the in vivo sensor before it may be introduced into a body. For example, an in vivo image sensor may be configured to capture images at fixed intervals beginning with the time that it is introduced into the
30 body. Typically, an in vivo sensor may be activated by a doctor or medical practitioner who assists in introducing such sensor into the body. Other in vivo sensors may be activated before ingestion, for example, automatically upon their removal from their original packaging. As a result, an in vivo sensor introduced to a location in the body that may be remote from an area of interest or suspected pathology in a body, may
35 perform its sensing functions or other functions in locations other than the area of interests for example where no pathology or suspected pathology exists. The

performance of such superfluous sensing may inefficiently utilize the power supply, data collection, data transfer (bandwidth), data storage capacity and/or other of the sometimes limited resource of the in vivo sensor. Redundant data may be required to be reviewed by the physician, increasing the overall review time.

5 The capturing of data by an in vivo sensor based on a fixed schedule may result on the one hand, in superfluous data being collected in areas that may be of little diagnostic or other interest, and, on the other hand, in insufficient sensory data being captured of in vivo areas that may be of particular diagnostic or other interest. For example, an in vivo image capturing system may be programmed to capture in vivo
10 images at a rate of, for example, two frames per second. While such frame capture rate may be for example sufficient to generally capture adequate images of most of the small bowel, such frame capture rate may be too slow to achieve the level of imaging detail that may be required for areas such as the esophagus or other areas.

 There is therefore a need for a system and method for allowing an efficient and
15 effective operation of an in vivo device.

SUMMARY OF THE INVENTION

 There is thus provided according to one embodiment of the invention, a system for
20 in vivo sensing including for example an in vivo sensing device with a condition tester, and a controller. The condition sensor may for example be operatively linked with the controller so as to control for example an operational mode of the in vivo sensing device.

 It is also provided according to an embodiment of the invention, a method for controlling, for example an in vivo imaging device by, for example, sensing a
25 condition in vivo and triggering an event in the in vivo imaging device based on the sensing.

BRIEF DESCRIPTION OF THE DRAWINGS

5 The present invention will be understood and appreciated more fully from the following detailed description taken in conjunction with the appended drawings in which:

 Figure 1A is a schematic illustration of an in vivo device that may be used in accordance with an embodiment of the present invention;

10 Figure 1B is a schematic illustration of a receiver in accordance with an embodiment of the present invention;

 Figure 1C is a schematic illustration of a data processor in accordance with an embodiment of the present invention;

15 Figure 2 is a schematic illustration of an in vivo device with a condition sensitive, color-changing material in accordance with an embodiment of the present invention;

 Figure 3 is a schematic illustration of a device with two image sensors in accordance with an embodiment of the present invention;

20 Figure 4 is a schematic illustration of a condition tester in the form of a coating in accordance with an embodiment of the present invention;

 Figures 5A and 5B are schematic illustrations of a floatable device according to an embodiment of the invention;

 Figure 6 sets forth a flow chart of the operation of a controller in accordance with an embodiment of the present invention;

25 Figure 7 sets forth a flow chart of the operation of a device in accordance with an embodiment of the present invention;

 Figure 8 sets forth a schematic diagram of a temperature triggered circuit in accordance with an embodiment of the current invention;

30 Figure 9 is a chart depicting a change in mode based on a pH trigger in accordance with an embodiment of the current invention; and

 Figure 10 is a chart depicting a change in mode initiated by a pH trigger and combined with a timed delay in accordance with an embodiment of the current invention.

DETAILED DESCRIPTION OF THE INVENTION

5 In the following description, various aspects of the present invention will be described. For purposes of explanation, specific configurations and details are set forth in order to provide a thorough understanding of the present invention. However, it will also be appreciated by one skilled in the art that the present invention may be practiced without the specific details presented herein. Furthermore, well-known features may be
10 omitted or simplified in order not to obscure the present invention.

 According to some embodiments of the invention, a system, method and device are provided for triggering an event such as, for example, activating or altering the operational mode of an in vivo device and/or a receiving (and/or processing, and/or reviewing) unit, typically located outside a patient's body, in response to in vivo
15 conditions as may be detected by an in vivo condition tester. Such activating, deactivating or altering operational modes may include for example, activating or deactivating one or more components of the in vivo device and/or the receiving unit, increasing or decreasing the power consumption, increasing or decreasing the level of illumination, increasing or decreasing the rate of sensing, such as, for example,
20 increasing the data capture rate from, for example, 2 images per second to for example, 14 images per second, or altering the sensing parameters such as, for example, in the case of an in vivo image sensor, increasing or decreasing the illumination intensity of the light sources or altering the image plane of the image sensor. Other operational modes may be changed and other data capture rates may be used. In certain embodiments,
25 more than one in vivo sensor may be included in a single device. A change in the operational mode of the device may in such embodiments include activating or deactivating one or both of such sensors or alternating the activation of such more than one sensor. For example, an in vivo image sensor may include two image sensors. A change in operational mode may in such example mean activating or deactivating one or
30 both of such image sensors, or alternating the activation of such image sensors. In other embodiments, one in vivo device may activate or deactivate one or more components in second in vivo device. Communication between two or more in vivo devices may be for

example through one or more external receivers or may be through for example direct communication between one or more in vivo devices.

In certain embodiments changes in the operational mode may for example include changes in the methods or procedures of processing sensory data obtained, and optionally transmitted, from the in vivo device. For example, sensory data such as images or ultrasound readings from endo-luminal areas that have villi may return distorted images as a result of the irregular surfaces of the villi. In certain cases, such distortions may be corrected through changes in the methods of processing of the sensory data by the data processor. For example, specific image processing algorithms may be activated. According to one embodiment methods of processing sensory data may be executed, for example, in an external receiving unit. In another embodiment the change may be in the mode of the data presentation (reviewing mode), e.g. presentation of the images in double image vs. single image mode.

The invention according to certain embodiments, comprises an in vivo device such as, for example, an in vivo image capture system, an in vivo condition tester such as, for example, any of an in vivo pH tester, blood detector, thermometer, pressure tester, spectral analytic image sensor, biosensor for biosensing, accelerometer, or motion detector, and a controller for linking the condition tester with the in vivo device and for signaling the change to be made in the operational mode of the in vivo device. Other condition testers may also be used as well as a combination of two or more condition sensor may be used. In one exemplary embodiment a biosensor may be used to sense, for example, colon specific flora in a colon. In another exemplary embodiment a pressure tester may be used to sense, for example, a change in pressure, such as a change in pressure pattern. For example, a drop in pressure may be sensed by a pressure tester, for example, when the device moves from the small intestine to the cecum (at the beginning of the colon). Various signals emitted by the condition tester such as mechanical, electrical, electromagnetic, chemical, or optical signals may also be used.

Embodiments of the present invention may be used with in vivo devices and recording/receiving and display systems such as various embodiments described in US Patent No. 5,604,531, assigned to the common assignee of the present application and incorporated herein by reference, and/or Publication Number WO 01/65995, also assigned

to the common assignee of the present application and incorporated herein by reference. Other in vivo systems, having other configurations, may be used.

Embodiments of the device may be typically autonomous and typically self-contained. For example, the device may be a capsule or other units where all the components are substantially contained within a container or shell, and where the device does not require any wires or cables to, for example, receive power or transmit information. The device may communicate with an external receiving and display system to provide display of data, control, or other functions. For example, power may be provided by an internal battery or a wireless receiving system. Other embodiments may have other configurations and capabilities. For example, components may be distributed over multiple sites or units. Control information may be received from an external source.

An in vivo imaging system for example, that may be included in an ingestible device such as a capsule may capture and transmit images of the GI tract while the capsule may pass through the GI lumen. In addition to the imaging system, a device such as a capsule may include, for example, an optical system for imaging an area of interest onto the imaging system and a transmitter for transmitting the image output of the image sensor. A capsule may pass through the digestive tract and operate as an autonomous video endoscope. It may image difficult to reach areas of the GI tract, such as the small intestine. Other devices may be included, and devices including sensors other than image sensors may be used. Configurations other than capsules may also be used.

Reference is made to Fig. 1A, a schematic illustration of an in vivo device in the form of for example, a swallowable capsule that may be used in accordance with an embodiment of the present invention. Device 40 may comprise an image sensor 46, an in vivo optical system 41 for focusing light reflected back from in vivo areas (not shown) onto image sensor 46, an illumination source 42, such as one or more light emitting diodes (LEDs) or other suitable sources, a dome 44 that may be useful, inter alia, for protecting the optical system from body fluids, a circuit or controller 48 for controlling the operational mode, such as for example settings of the device 40, a condition tester 49 such as for example, a pH tester or thermometer, an in vivo memory unit 39, an in vivo power source 45 such as a set of batteries, an in vivo receiver 43 for collecting signals transmitted to device 40, and an in vivo transmitter 47 for transmitting signals and/or

image data to a receiver. One or more of in vivo image sensor 46, in vivo illumination source 42, controller 48, in vivo memory unit 39, in vivo transmitter 47, in vivo receiver 43 and condition tester 49 may in certain embodiments of the present invention be operatively connected, for example to/or through PCB 38, or included or embedded within an application specific integrated circuit (ASIC) 50. In other embodiments, image sensor 46, controller 48 and condition tester 49 may be operatively linked to each other without an ASIC 50 or PCB 38 or other connecting means. A wired or wireless connection, such as for example a microwave connection or other suitable connections may be used between elements in the capsule. Such an ASIC 50 may provide control for the capsule. Alternatively, another component such as transmitter 47 may provide such control.

In certain embodiments, image sensor 46 may be a CCD or a CMOS image sensor that may have arrays of various typically color pixels. Other suitable image sensors or no image sensors may be used. In one embodiment of the invention, image sensor 46 may also function as a condition tester. For example, an image sensor may be used to detect for example, blood vessel structures typically found in colon, or villi structures typically found in small intestine. Detection of such structures, detection of lack of such structures, or detection of other structures or colors such as for example color specific to content in the intestine may be used to trigger an event in the in vivo device. Other suitable structures or colors detected may be used as a trigger. Detection, according to an embodiment of the invention, could be aided by appropriate image processing algorithms and/or suitable software.

In other configurations of device 40, components such as capsule receiver 43, power source 45, in vivo memory unit 39 or other units may be omitted.

Typically, device 40 is swallowed by a patient and traverses a patient's GI tract. Other suitable body lumens or cavities may be imaged or examined.

Reference is now made to Fig. 1B, a schematic illustration of an external receiver 12 in accordance with an embodiment of the present invention. External receiver 12 may typically be located outside the patient's body and may receive and/or record and/or process the data transmitted from device 40. External receiver 12 may typically include a receiver antenna (or antenna array) 15, for receiving image and other data from device 40 and stored in for example storage unit 16. Typically, external receiver 12 may be portable, and may be worn on the patient's body during recording of the images.

External receiver 12 may also be equipped with processing unit 11, such as for example signal processing unit and/or control software or for example a control mechanism or circuit emulating such functionality that may control for example, evaluate and respond to signals transmitted by device 40. External receiver 12 may also include a transmitter and receiver transmitter 17 that may enable external receiver 12 to transmit signals such as control signals to device 40. External receiver 12 may also include a user interface (not shown) that may inter alia provide indications to a user or patient as to changes made in the operational mode of a device. For example, passage of a capsule through the stomach may be identified by changes detected in pH levels that may for example trigger a change in the operational mode of a sensor such as an image sensor. A patient may be signaled via a user interface that such mode change is being made and prompted to take certain actions such as for example, changing positions (such as for example, changing from a sitting position to a reclining position), ingesting a laxative, or certain liquids, etc.

Reference is now made to Fig. 1C, a schematic illustration of a data processor in accordance with an embodiment of the present invention. Preferably, data processor 14, data processor storage unit 19 and monitor 18 are part of a personal computer or workstation that may include standard components such as a processor 13, a memory (such as storage unit 19, or other memory), a disk drive, and input-output devices. Alternate configurations are possible. In alternate embodiments, the data reception and storage components may be of another configuration. Further, image and other data may be received in other manners, by other sets of components. Typically, in operation, image data is transferred from external receiver 12 to data processor 14, which, in conjunction with processor 13, storage 19, and software, stores, possibly processes, and displays the image data on monitor 18. Other systems and methods of storing and/or displaying collected image data may be used. In other embodiments, processing of data can be performed by components within the external receiver 12.

Typically, device 40 may capture an image and transmit the image by using, for example, radio frequencies, to receiver antenna(s) 15. In alternate embodiments external receiver 12 is an integral part of data processor 14. Typically, the image data recorded and transmitted is digital color image data, although in alternate embodiments other suitable image formats (e.g., black and white image data, infrared image data, etc.) may be used. In one embodiment, each frame of image data may include 256 rows of 256 pixels each, each

pixel including data for color and brightness, according to known methods. For example, color may be represented in each pixel by a mosaic of four sub-pixels, each sub-pixel corresponding to primaries such as red, green, or blue (where one primary is represented twice). The brightness of each sub-pixel may be recorded by, for example, a one byte (*i.e.*,
5 0-255) brightness value. Other data suitable formats may be used. In one embodiment, image sensor 46 may capture or transmitter 47 may transmit image or other data in a diluted mode, capturing or transmitting for example, 16 rows of 16 pixels each.

In an embodiment, in vivo transmitter 47 may include at least a modulator (not shown) for modulating the image signal from the image sensor 46, a radio frequency (RF)
10 amplifier (not shown), and an impedance matcher (not shown). The modulator may convert the input image signal that may have for example, a cutoff frequency f_c of less than 5 MHz to an RF signal having a carrier frequency f_r , that may typically be in the range of 1 GHz. The carrier frequency may be in other bands, *e.g.* a 400MHz band. The modulated RF signal may typically have an appropriate bandwidth of f_b . The impedance matcher may
15 match the impedance of the circuit to that of the antenna. Other suitable transmitters or arrangements of transmitter components may be used, utilizing different signal formats and frequency ranges. In one embodiment of device 40, transmission may occur at a frequency for example of 434MHz, using for example Phase Shift Keying (PSK) or MSK (Minimal Shift Keying). In alternate embodiments, other suitable transmission frequencies and
20 methods, such as for example AM or FM may be used.

External receiver 12 may detect a signal having the carrier frequency f_r and the bandwidth f_c such as described hereinabove. External receiver 12 may be similar to those found in televisions or it may be one similar to those described on pages 244-245 of the book "Biomedical Telemetry" by R. Stewart McKay and published by John Wiley and
25 Sons, 1970. The receiver may be digital or analog. In alternate embodiments, other receivers, responding to other types of signals, may be used.

In certain embodiments, condition tester 49 may be an in vivo pH tester, as is well known in the art, for example a pH tester using the technology used in known pH measuring capsules. Such pH tester may utilize as electrodes an external ring electrode
30 made of antimony and the zinc-silver chloride electrode of the battery that powers the tester. A saline solution such as for example, a 0.9% physiologic saline solution may be introduced into the electrode chamber immediately prior to the testing. The potential

difference that develops between the two electrodes and that depends on the pH may be applied to a transistor as a frequency-determining measuring voltage.

Other pH testers, such as ion selective field effect transistors (ISFET), may also be used as condition tester 49 to evaluate pH in areas adjacent to the location of the device 40. ISFET sensor chips that may be used for in vivo pH detection are known in the art as may be described, for example, in Wang, L., Integrated Micro-Instrumentation for Dynamic Monitoring of the Gastro-Intestinal Tract, as presented at the IEEE Instrumentation and Measurement Technology Conference, May 2002, retrieved on October 15, 2002 from the Internet: <URL: <http://www.see.ac.uk/naa.publications.html>>.

Other suitable pH testers may also be used. An ISFET sensor serving as condition tester 49 may be operatively connected to ASIC 50 or otherwise may be connected directly to image sensor 46. In a typical embodiment, an ISFET sensor serving as condition tester 49 may be situated adjacent to the outer wall of the device 40 so as to maximize the exposure of such condition tester 49 to the in vivo conditions outside of such wall of device 40.

In some embodiments, controller 48 may be substituted or complimented by an external controller located out of the body. For example the external controller may be an integral part of processor 11. In such embodiments, triggering may be external triggering. Condition tester 49 may transmit a signal to in vivo transmitter 47 that transmits such signals to receiver antenna(s) 15. External receiver 12 may process such signals and transmit back triggering signal such as instructions by way of receiver transmitter 17 to in vivo receiver 43. In vivo receiver 43 may then direct a change in the mode of operation of device 40. In some embodiments, external receiver 12 may be capable of overriding or initiating a change in the mode of operation of device 40 in response to a signal that is input to receiver by medical personnel.

A condition tester such as for example, a pressure sensor may use a strain gauge as a condition detector, such as for example, a thin foil, typically a semiconductor or a piezoelectric material. Such strain gauge may accept power through a wire and provide a variable strain signal on such wire.

In other embodiments, condition tester 49 may take the form of a condition sensitive, color-changing material. Reference is now made to Fig. 2, which is a schematic illustration of an in vivo sensor with a condition sensitive, color-changing material 202 in accordance with an embodiment of the present invention. Material 202

may be temperature sensitive. "Temperature-sensitive" in the context of the present invention may be defined as reactive to a change in temperature. This temperature change may include a range of temperatures or a change from a reference temperature to another temperature. In other embodiments, material 202 may be pressure-sensitive, pH sensitive or sensitive to the presence of certain substances such as for example, blood, with color-changing characteristics varying with changes in such conditions. Thus, different properties within the environment of the body lumen can be measured in a manner similar to the one described for temperature hereinbelow. For example, a pH condition tester may use litmus paper as a color-changing material 202, and a blood detector may use a polyelectrolyte as a color-changing material 202, as known in the art.

In an embodiment, the temperature-sensitive, color-changing material 202 may be a Thermotropic Liquid Crystal (TLC) paint or coating, such as are offered by Hallcrest, Inc. of Glenview, Illinois. The TLCs, that may, for example, be cholesteric (including sterol-derived chemicals) or chiral nematic (including non-sterol based chemicals) liquid crystals, or a combination of the two, provide color changes in response to temperature changes. These color changes may be reversible or hysteretic. In certain embodiments that include materials 202 that may be capable of reversible color changes, controller 48 may be programmed to reverse or further alter the operational mode changes in image sensor 46 in the event that a condition tester ceases to detect the changed color of material 202.

The TLC can be used in several forms according to several embodiments, including but not limited to paints, microencapsulated coatings and slurries, TLC coated polyester sheets, and unsealed films.

As shown in Fig. 2, temperature-sensitive color-changing material 202 may be placed on the inside of capsule 200, with color-changing portions facing inwards. By placing material 202 on the inside of the capsule, potential problems associated with the biocompatibility and the resilience of material 202 in light of bodily fluids and pH changes may be avoided. However, it should be apparent that color-changing material may also be placed on the outside of capsule 200 where it may be in contact with bodily fluids. Such contact between material 202 and bodily fluids may facilitate testing of such bodily fluids for reactions with material 202. In certain embodiments, it may be

necessary to achieve contact between bodily fluids and material 202. The attachment or placement of material 202 can be accomplished in several ways. For example, material 202 may be in the form of paint, and may be painted onto the capsule. In another embodiment, material 202 may be attached onto the capsule with adhesive. In a further embodiment, material 202 may be sprayed onto the dome 44 as a coating. In yet a further embodiment, material 202 may be enclosed in a semi-permeable membrane in contact with bodily fluids.

In the course of the function of capsule 200, light from a light source 204 may be directed towards material 202. Light source 204 may include one or several components, preferably light emitting diodes (LEDs) that may be placed in various locations within capsule 200. Light source 204 may also be used as or provided by illumination source 42 shown in Fig. 1, to illuminate the environment being imaged (outside of the capsule), or a separate illumination source 204 may be included for that purpose.

Changes in in vivo conditions, such as, for example, changes in temperature, pH, pressure, the presence of blood and the like (depending on the nature of material 202), may in certain embodiments cause various materials that can be used as color-changing material 202, to change color. Image sensor 46 detects the appearance of the new color when light from light source 204 is reflected back from material 202 onto image sensor 46. Referring to Fig. 2, such detection of changes in color may in certain embodiments be performed by a subgroup of pixels 206 included in the pixel array of image sensor 46. In one embodiment of the invention, pixel array of image sensor may have one subgroup of pixels that are sensitive to a first range of wavelengths e.g., colors and another subgroup of pixels sensitive to second range of wavelengths, e.g., colors. In some embodiments such one subgroup of pixels, or specific pixels may be positioned on the pixel array of image sensor 46 to be exposed to light reflected back from material 202 considering the angle of incidence 208 and angle or return 208' of the light directed onto and reflected back from material 202. Similarly, in certain embodiments, such subgroup of pixels 206 may be sensitive to a specified range of colors that appear on material 202 once the designated in vivo environmental condition may be detected. In an alternative embodiment special photodiode(s) may be used in addition to or in place of a subgroup of pixels 206 to detect color changes.

When a designated change in color of material 202 is detected by a subgroup of pixels 206, a signal may be sent to controller 48 by such a subgroup of pixels 206 or by another component operatively connected to a subgroup of pixels 206. In certain embodiments, a subgroup of pixels 206 may be replaced or supplemented by a spectral analyzer that is capable of detecting color changes in material 202. Other color-sensitive detectors may also be used. Such detection or processing may also be aided or performed by a processor or circuitry located in ASIC 50, external receiver 12 or data processor 14.

In certain embodiments, a range of color sensitive pixels, some of which may be sensitive to the various colors that can appear on material 202 may be situated on pixel array 210 of image sensor 46. Signals produced by each of such specific pixels 206 may vary depending on the color appearing on material 202. Controller 48 may detect and differentiate between such various signals, for example by utilizing appropriate image processing algorithms, and issue instructions to a sensor in response to each thereof. According to one embodiment a change of color may be detected in the in vivo environment that is being imaged. For example, a spot of bleeding may appear in a certain image. The change of color, that may indicate, for example, pathology in the GI tract, may be recognized by known methods. For example, controller 48 or data processor 14 may generate a probability indication of presence of colorimetric abnormalities on comparison of color content of the images and at least one reference value, for example, as described in PCT publication WO 02/073507, published on 19 September, 2002, that is assigned to the common assignee of the present invention. According to some embodiments, once a color change may be detected the controller 48 or data processor 14 may initiate a change in the mode of operation of device 40, of the external receiver 12, of both or of any other component or combination of components of the system. In other embodiments, a photodiode maybe used to detect changes in material 202. Such photodiode may in certain embodiments be connected to an amplifier that may be further connected to a comparator. A mode change may thereby be triggered by analog rather than digital electronics.

In one embodiment of the invention, one or more photodiodes may be used to detect light, such as for example, visible light, IR light, or other ranges of light illuminated for example externally through the skin toward an in vivo area of interest. A

photodiode or other light detecting unit, for example incorporated in an in vivo device may sense illumination when approaching for example toward such area of interest. Such detection may trigger a change in operational mode. Other suitable signals besides light may be used to penetrate the skin or other tissue and other suitable detection units
5 may be used to pick up penetrated signal in vivo. For example, an acoustic signal may be used.

In an embodiment, capsule 200 may operate in a low power consumption mode until a color change in material 202 may be detected. For example, until such color change may be detected, light sources 204 may be set to illuminate once every second,
10 thereby consuming less power than used by the overall capsule 200 during full operation that might in certain embodiments illuminate several times a second or more. In response to a signal that may be detected from specific pixels 206, controller 48 (or another component located in capsule 200, external receiver 12 or data processor 14) may alter the mode of operation of capsule 200 or of any other component of the system. For example,
15 in certain embodiments, any or both of light source 204 and image sensor 46 may be directed to increase the rate of capture of images in order to more fully image the endoluminal vicinity wherein a specific condition may have been detected. Controller 48 may direct other activations or alterations in the mode or operation of capsule 200. In other embodiments, the response of controller 48 to signals from specific pixels 206, may be,
20 for example, any of turning on the image sensor 46 that may theretofore have been inactive, changing mode of image sensor or transmitter, collecting samples of in vivo liquids or other materials, releasing encapsulated drugs that were held in capsule 200 or performing other functions.

In some embodiments, the pixels receiving the color indication may be, for
25 example, the regular pixels of image sensor 46. Post processing circuitry or software located in capsule 200, external receiver 12 or data processor 14 may analyze the signals from the set of pixels (set being understood to include one unit) and make a mode change determination therefrom.

Other embodiments besides colorimetric changes may include, for example,
30 temperature measurement using devices such as thermistors (located in a capsule for example as a discrete component or as part of ASIC) or using pH electrodes, and other embodiments.

Reference is now made to Fig. 3 that is a schematic illustration of a capsule 300 with two image sensors in accordance with an embodiment of the present invention. Capsule 300 has one image sensor 302 at one end of capsule 300 and a second image sensor 304 at another end of capsule 300. In an embodiment of the present invention, condition tester such as a color-changing material 202 such as those described in Fig. 2 may be installed proximate to image sensor 302, and such image sensor 302 may in such embodiment have specific pixels 206 similar to those described above for detecting color changes in material 202. When a change in color of material 202 is detected by image sensor 302, a signal of such change is sent to controller 48 of capsule 300. Controller 48 may in such embodiment alter the operational mode, such as for example by activating a component, for example the image sensor 304 of capsule 300. For example, the operational mode of both or either of image sensors 302 and 304 may be changed. Such a mode change may, for example, increase the number of images to be captured of such area or alter the orientation of images captured or differential activation of either one or both image sensors may be affected in response to a signal, or other mode changes discussed herein.

In certain embodiments, controller 48 may be configured to delay issuing operational mode change orders to until more than one signal from condition detector 49 may have been received. In an embodiment of the present invention, controller 48 may be configured with a delay mechanism in the form of for example a counter 51 that causes controller 48 to delay activating or altering the operational mode of image sensor 304 until several signals from condition tester 202 may have been received, or until signals signifying that a certain condition exists may be received over the course of a certain period of time. Such activation may, for example, reduce the chance that a false reading or fleeting condition activates image sensor 304, or may provide "debouncing" in case conditions may change in a variable manner between one relatively steady state and another. For example, in one embodiment, capsule 300 may operate in a first mode (e.g., low power consumption, or at a first frame capture rate) in the mouth and esophagus, where the pH is generally approximately 7-8. When capsule 300 reaches the stomach, where the pH is typically about 2, a pH detector on or within capsule 300 may detect a change in pH, and the operational mode may change, for example to a different power consumption, or a different frame capture rate. Later, when capsule 300 reaches

the small intestine, capsule 300 may detect a change in pH to, for example 7-8, and the operational mode may change again. A change in pH may cause alteration in the operational mode only if received for, for example, one minute (other suitable time periods may be used). Other methods of debouncing or guarding against fleeting
5 conditions may be used.

Controller 48 may in certain embodiments be a software controller embedded into ASIC 50. In other embodiments, controller 48 may be a simple switch or circuit connected to for example a condition tester such as a thermistor 800. The controller may include, for example, an amplifier 802 and a comparator 804, comparing the measured
10 signal to some pre-defined threshold 806, as are depicted forth, for example, in Fig. 8. Such switch or circuit may in certain embodiments power on or trigger the activation of ASIC 50 when the proper condition may be detected. In other embodiments, such switch or circuit may signal ASIC 50 to, for example, begin operation or change the mode of operation of the sensor.

15 In a further embodiment, the switch from one condition and then back to another may be the trigger for a mode change. For example, in case a high pH is detected for a period, then a low pH, then again a high pH, the mode change may occur only on the third condition change. Other suitable signals or series of signals may be used to trigger other suitable functionalities. Further, altering the mode based on detection of a
20 condition change may be combined with, for example, a delay. For example, capsule 300 may wait, for example, one hour after detecting a condition change to effect a mode change. Fig. 4 is a schematic illustration of a condition tester in the form of a coating in accordance with an embodiment of the present invention. In such embodiment, a portion of capsule 400 may be coated with one or more layers of a dissolvable material
25 402. Each layer of dissolvable material 402 may be comprised of varying substances that dissolve at varying rates or when exposed to specific materials or environments. For example, a first, outer layer 404 of dissolvable material may be pH sensitive and dissolve when exposed to the acidic environment of the stomach, and may expose certain components such as for example switches 412, sensors 408 or drug compartment
30 410 with an opening, while capsule 400 may be in a specified site such as for example, the stomach. A second inner layer 406 may for example, dissolve in the more basic environment of the small intestine and may activate other sensors or release other

encapsulated drugs. Other materials that may be sensitive to elapsed time and dissolve in accordance with a specific period of time after introduction to the GI tract may also be possible as a means of delaying activation of certain functions of capsule 400. An example of dissolvable materials that may be used as such coatings include starches, such as gelatinous materials, waxes, biodegradable plastics, and other known biodegradable materials. Other suitable dissolvable materials with other characteristics may also be used.

Dissolvable material 402 may cover any or all of a sensor 408, such as for example, a pH sensor, a switch 412, such as for example a switch that turns on an image sensor, an encapsulated drug compartment 410 that releases its contents or a sampling inlet 414 that lets surrounding fluids enter a compartment where such fluids may be sampled, captured or evaluated by a sensor. When dissolvable material 402 dissolves, sensor 408 may be exposed, switch 412 may be activated, sampling inlet 414 may be opened or an encapsulated drug compartment 410 may release its contents into the surrounding area. In other embodiments, the dissolving of dissolvable material 402 may facilitate contact between electrical leads that had theretofore been separated, such contact may signal a change in operational mode. According to another embodiment a magnet may be held in the vicinity of the capsule 400 such that it affects the ON/OFF status of the capsule. In some embodiments the magnet may be embedded in a dissolvable coating, such as dissolvable material 402, such that while the coating is intact, the capsule is OFF. When the coating dissolves, for example, in response to environmental pH, the magnet may be freed and may become dissociated from the capsule allowing the capsule to be ON. In other embodiments other suitable environmental triggers may cause the dissolving of coatings.

Another embodiment is schematically illustrated in Figs. 5A and 5B. In this embodiment an imaging capsule 500 may be a floatable capsule, for example, a capsule having a specific gravity of less than 1. A floatable capsule is described, for example, in Publication Number WO 02/095351, published on November 28, 2002 assigned to the common assignee of the present inventions and is hereby incorporated in its entirety by reference. Such a capsule may be advantageous for passage through portions of a voluminous cavity, such as the stomach and/or large intestine. In other portions of voluminous cavities (e.g., the descending portion of the large intestine) a floatable

capsule may be delayed rather than advanced. Thus, a floatable capsule may benefit from having the option of loosing its floatation characteristics at a given point during its passage through the GI tract, for example, while in the large intestine.

According to one embodiment, a capsule may have a fluid chamber such as for
5 example a floatation compartment 502 that may be filled with a fluid, a gas, or other suitable material that is lighter than the endo-luminal fluid, for example, air. In certain embodiments, floatation compartment may be as small as 5% of the volume of capsule 500. Other suitable volumes may be used. The floatation compartment 502 may have a valve 504 keeping the compartment 502 closed and the capsule 500 floating. Upon
10 triggering, valve 504 may be opened (see Fig. 5B). Floatation compartment 502 may then be filled with endo-luminal liquid, raising the specific gravity of capsule 500 and rendering capsule 500 non-floating. As such the floatation mode of a capsule may be altered.

A number of mechanisms for opening valve 504 may be implemented, such as,
15 electronic, mechanical or chemically based mechanisms. For example instant heating (requiring only a small amount of battery energy) may be applied, melting material of valve 504. The signal for effecting the change may be as described above.

Fig. 6 sets forth a flow chart of the operation of a controller 48 in accordance with an embodiment of the present invention. Such controller may in an embodiment be
20 a software controller in the form of logic programmed into, for example ASIC 50, controller 48, external receiver 12 or other suitable components. Such software controller may have a flag to indicate the operational mode to which a sensor is set. Settings of such flag may be 0 or 1 for on or off, or other suitable settings to indicate other settings to which a sensor may then be operating. Software controller may also
25 include a counter that may in certain embodiments count signals received from condition tester 49 indicating the detection of the conditions to be tested by condition tester 49. Software controller may also be operatively linked to an operation activator of an in vivo component such as image sensor 46 that controls the operation of such sensor. For example, operation activator may be an internal clock that controls the timing of the
30 image capture rate of image sensor 46 or the operation of light source 204.

In its initial state 602, the flag of software controller may be set to 0, the counter may be set to 0 and the activator may be set to 0. In such settings, image sensor 46 may

not be capturing images or may be in some other reduced mode of operation. In step 604, condition tester 49 may detect a changed condition in the in vivo area surrounding capsule 40 and may signal software controller 48 as to such changed condition. Such signal increments counter to 1. Step 604 may be repeated by condition tester 49 at periodic intervals that match the sampling rate of condition tester 49. Each signal delivered by condition tester 49 that indicates the changed condition may increment the counter by 1 (605). Once the counter reaches a designated threshold in step 606, the flag switches to 1 in step 608. Such switch by the flag to 1 switches the sensor activator to 1 as in Step 610. The activator may then change the mode of operation of image sensor 46. Such change may for example be an increase in the frame capture rate of image sensor 46 or any other suitable change in the operational mode of the sensor.

In certain embodiments, the counter may be decremented each time condition tester 49 sends a signal to controller 48 that indicates the absence of an elevated condition, thereby possibly indicating that conditions may have returned to pre-defined normal levels. Once the counter may have been decremented below a pre-defined threshold level, the flag may revert to 0 and may reset the activator to its initial setting so that such sensor may resume the operational mode that was in effect prior to the change described above, or some other suitable operational mode.

In other embodiments, condition tester 49 may be, for example, a clock such as for example an internal clock embedded into ASIC 50 or otherwise operatively connected to image sensor 46. In such case, controller 48 may be a component such as for example a switch operatively attached to such embedded clock that may turn on once a designated period has elapsed. Such elapsed period may be the estimated time that it takes capsule 300 to pass through the stomach and into the small intestine where the desired image capturing may take place. Other periods may also be designated depending on where in the GI tract the desired image capturing may be designated to begin.

In yet further embodiments, an ingestible capsule may be meant for imaging or otherwise sensing distal portions of the GI tract, such as the large intestine. A method for economically using an imaging (or other sensing) capsule is provided according to an embodiment of the invention. Figure 7 illustrates a method for imaging or otherwise

sensing distal parts of the GI tract according to an embodiment of the invention. An inactive device such as for example a capsule (e.g., does not sample or transmit images or other data) is swallowed (710) by a patient. According to one embodiment the capsule may comprise temperature sensing capabilities. Any in vivo temperature sensing
5 mechanism, such as those known in the art, may be used. After a capsule is swallowed a patient may be made to ingest a volume of cold or hot water (720) at regular intervals. According to one embodiment the patient may ingest cold or hot water over a period of a few hours (e.g., 3-5 hours), for example, a period in which the capsule has most probably left the stomach. According to another embodiment the patient may be made to
10 ingest a volume of cold or hot water until alerted that the capsule has left the stomach (further detailed below). While the capsule may be in the stomach an ingested volume of cold or hot water may cause a change of temperature in the stomach environment. Once in the small intestine, the effect of a cold or hot drink may no longer be felt. According to one embodiment a capsule may be programmed to sense a periodical change in
15 temperature (700), for example to sense a temperature above or below a certain threshold, at predetermined intervals. While a temperature change may be sensed at predetermined intervals, the capsule may be kept inactive (701). If a temperature change is not sensed at one predetermined time, the capsule may be triggered (for example, as detailed above) to activate the image sensor or other components (702). Thus, the
20 capsule may begin collecting data only after leaving the stomach for example, such that it is closer to the large intestine thereby saving energy and allowing effective and complete action of the capsule in the large intestine.

According to some embodiments activating the capsule may cause a signal to be transmitted (703) to an external receiving unit so as to activate an alert 730 (e.g., a beep
25 or a flashing light), that may alert a patient to start or stop an action for example to stop drinking the cold or hot drink (740). Also, the patient may then be prepared for the expected imaging or otherwise sensing of the large intestine, for example, the patient may thus be warned to begin taking a laxative.

Reference is made to Fig. 9 that is a chart depicting a change in mode based on a
30 pH trigger in accordance with an embodiment of the current invention. As depicted in Fig. 9, a device may be in a first operational mode from, for example beginning with the time it is turned on and while it is for example, in the stomach wherein pH is low. As

the device may leave the stomach, pH may rise. Such rise may set off the pH trigger that may change the operational mode of the device. Other suitable triggers may be used as well. Such change may, for example, be a component such as for example a switch of the device imaging with two image sensors 302 and 304 (as are depicted, for example, in
5 Fig. 3) to imaging with only a first image sensor 302. In such an embodiment effective viewing of the upper regions of the GI tract may be enabled, by using two image sensors whereas, a power saving mode may then be switched to in the small intestine where one image sensor may be enough to provide effective viewing.

Reference is made to Fig. 10 that is a chart depicting a change in mode initiated
10 by a pH trigger and combined with a timed delay in accordance with an embodiment of the current invention. As depicted in Fig. 10, a device may be in a first mode of operation immediately when it is introduced into a body. The mode of operation may change, such as for example, going to off or some other inactive state, until a trigger occurs such as for example a change in pH. Other suitable triggers may be used as well.
15 The trigger may initiate, for example, a time delay during which the mode of operation may remain initially unchanged, but during which the device counts down until the delay ends, whereupon the mode change may be implemented. A trigger combined with a time delay may be useful for example where the large intestine may be the area to be imaged. In such an embodiment, the trigger may be the pH change that occurs when the
20 device leaves the stomach. The time delay may be the approximate time required for the device to traverse the small intestine (e.g., 3-6 hours). Once the device nears the large intestine it may change modes of operation to image the desired area. In this way, the device may preserve its power supply until many hours after it is introduced into a body and until it reaches the targeted imaging area. Other suitable combinations of time
25 delays and triggers are possible.

It will be appreciated by persons skilled in the art that the present invention is not limited by what has been particularly shown and described herein above. Rather the scope of the invention is defined by the claims.

We claim:

1. A system for in vivo sensing, said system comprising:
an in vivo sensing device, said device comprising a condition
tester; and
5 a controller to control an operational mode of said in vivo
sensing device;
wherein said condition tester is operatively linked with said
controller.
2. The system according to claim 1 comprising an image sensor.
- 10 3. The system according to claim 2 wherein the image sensor is selected from a
group consisting of: CCD and CMOS.
4. The system according to claim 2 wherein the image sensor comprises one
subgroup of pixels said one subgroup being sensitive to a first range of colors,
and another subgroup of pixels, said other subgroup of pixels being sensitive to
15 a second range of colors.
5. The system according to claim 4 comprising a spectral analyzer.
6. The system according to claim 1 wherein the condition tester is selected from a
group consisting of: a pH tester, a blood detector, a thermometer, a pressure
sensor, a biosensor, a spectral analytic image sensor, an image sensor, and a
20 counter.
7. The system according to claim 1 wherein the condition tester is to test in vivo
conditions.

8. The system according to claim 1 wherein the controller is incorporated in the in vivo sensing device.
9. The system according to claim 1 wherein the controller is an external controller.
- 5 10. The system according to claim 1 wherein the controller comprises a counter.
11. The system according to claim 1 wherein the controller is selected from a group consisting of: mechanical switch, software, and circuitry.
12. The system according to claim 1 wherein the controller is a circuit, said circuit comprising an amplifier and a comparator.
- 10 13. The system according to claim 12 wherein the condition tester is a thermistor.
14. The system according to claim 1 comprising an in vivo transmitter.
15. The system according to claim 1 comprising an in vivo illumination source.
16. The system according to claim 1 comprising a photodiode.
17. The system according to claim 1 wherein the in vivo sensing device is an
15 autonomous device.
18. The system according to claim 1 wherein the in vivo sensing device is a capsule.
19. The system according to claim 1 wherein the in vivo sensing device comprises an ASIC wherein said ASIC is operatively connected to a component of the in
20 vivo sensing device.
20. The system according to claim 19 wherein the component is selected from the group consisting of: an in vivo transmitter, an in vivo illumination source, an

in vivo power source, a controller, an in vivo image sensor, a condition tester, an in vivo receiver, and an ASIC wherein said ASIC is operatively connected to the in vivo receiver.

21. The system according to claim 19 wherein the controller is an integral part of the ASIC.
22. The system according to claim 1 comprising an in vivo receiver.
23. The system according to claim 1 comprising an external receiver.
24. The system according to claim 1 wherein said external receiver includes a processing unit and a storage unit.
25. The system according to claim 1 comprising a monitor and a data processor.
26. The system according to claim 25 wherein said data processor comprises a storage unit and a processor.
27. The system according to claim 1 wherein the condition tester includes a color-changing material.
28. The system according to claim 27 wherein the color-changing material is selected from a group including: temperature sensitive material, pH sensitive material, and a blood sensitive material.
29. The system according to claim 1 wherein the condition tester includes a layer of pH sensitive and/or time sensitive dissolvable material.
30. The system according to claim 1 wherein the in vivo sensing device comprises a compartment coated with a pH sensitive and/or time sensitive dissolvable material.

31. The system according to claim 1 wherein the in vivo sensing device comprises a sampling inlet coated with a pH sensitive and/or time sensitive dissolvable material.
- 5 32. The system according to claim 1 wherein the in vivo sensing device comprises a switch coated with a pH sensitive and/or time sensitive dissolvable material.
33. A method for controlling an in vivo imaging device said method comprising:
- sensing a condition in vivo; and
- triggering an event in said in vivo imaging device based on said
- 10 sensing.
34. The method according to claim 33 wherein sensing a condition in vivo is selected from a group consisting of: time sensing, pH sensing, temperature sensing, pressure sensing, blood sensing, and biosensing.
35. The method according to claim 33 wherein the triggering is by a controller.
- 15 36. The method according to claim 33 wherein the triggering is by an external receiver.
37. The method according to claim 33 wherein the event comprises a change in an operational mode of the in vivo imaging device.
38. The method according to claim 37 wherein the change in operational mode is
- 20 selected from a group consisting of: activating a sensor, deactivating a sensor, altering data capture rate; altering signal format and frequency range of transmission; altering processing of sensory data; altering frame capture rate of an in vivo image sensor, altering illumination intensity, altering image plane of

an in vivo image sensor, activating in vivo sample collection, releasing a drug, altering power consumption mode, and altering floatation mode.

39. The method according to claim 33 comprising delaying triggering of an event.

40. The method according to claim 33 comprising ingesting a volume of cold or
5 hot water.

41. The method according to claim 33 wherein the triggering is by a pH sensitive and/or time sensitive dissolvable material.

ABSTRACT

A device, system and method for selectively activating or altering the operational mode of an autonomous in vivo device in response to in vivo conditions. The system includes an in vivo sensing device with a condition tester, and a controller. The in vivo
5 sensing device may be in communication with an external receiver.

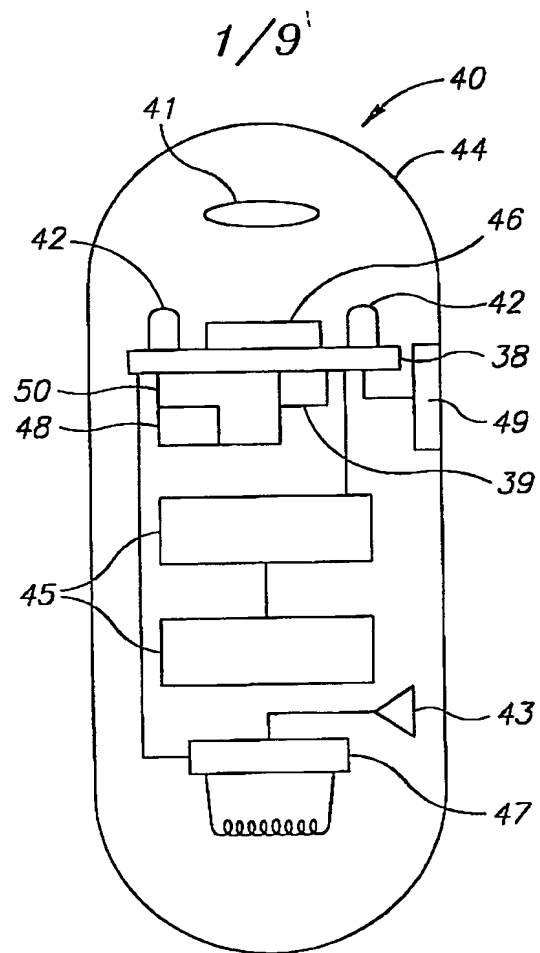


FIG.1A

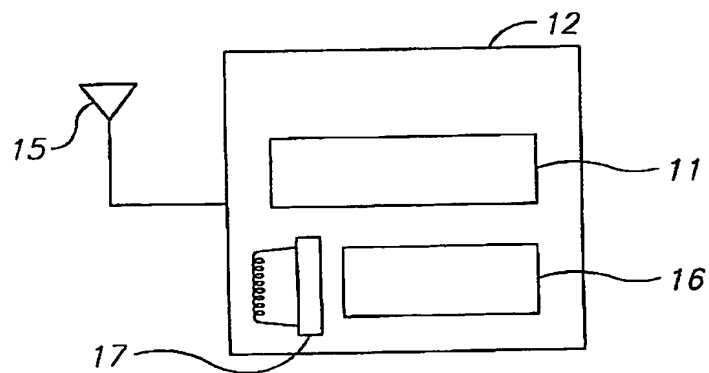


FIG.1B

2/9

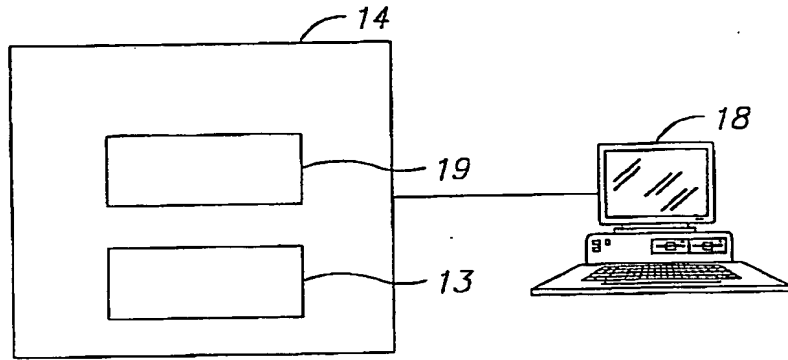


FIG. 1C

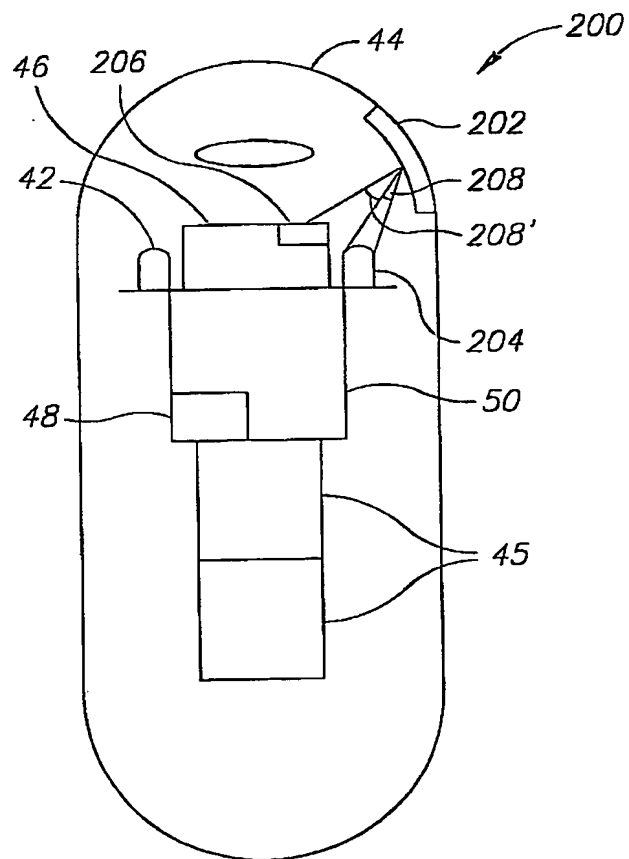


FIG. 2

3/9

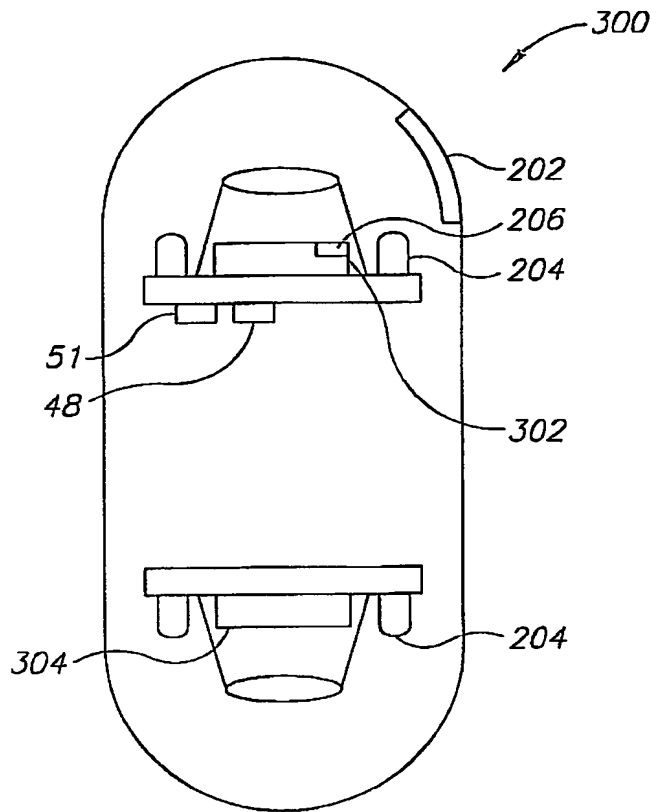


FIG. 3

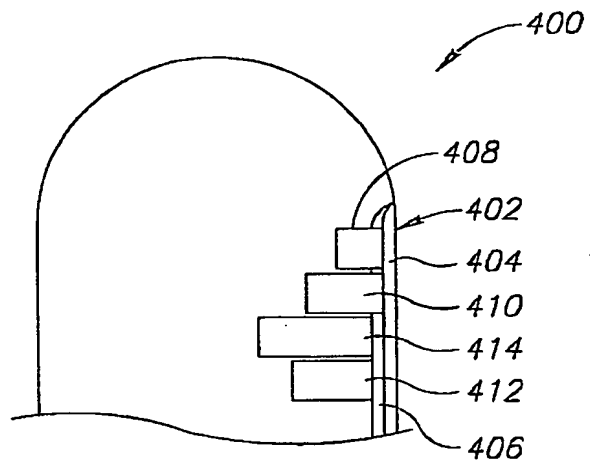


FIG. 4

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500

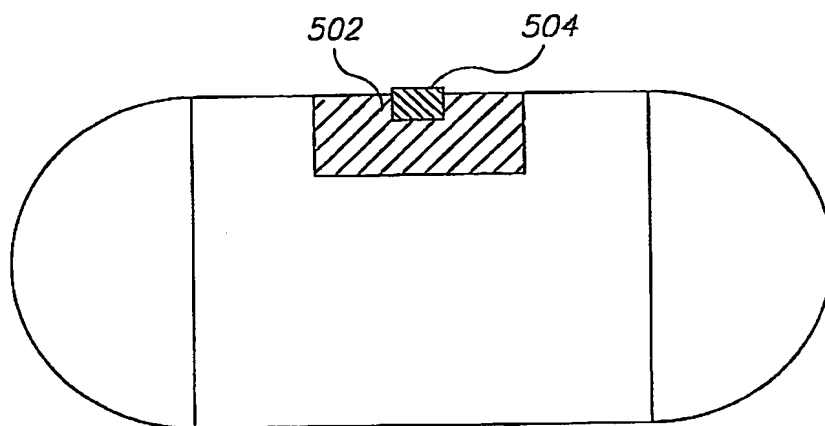


FIG. 5A

500

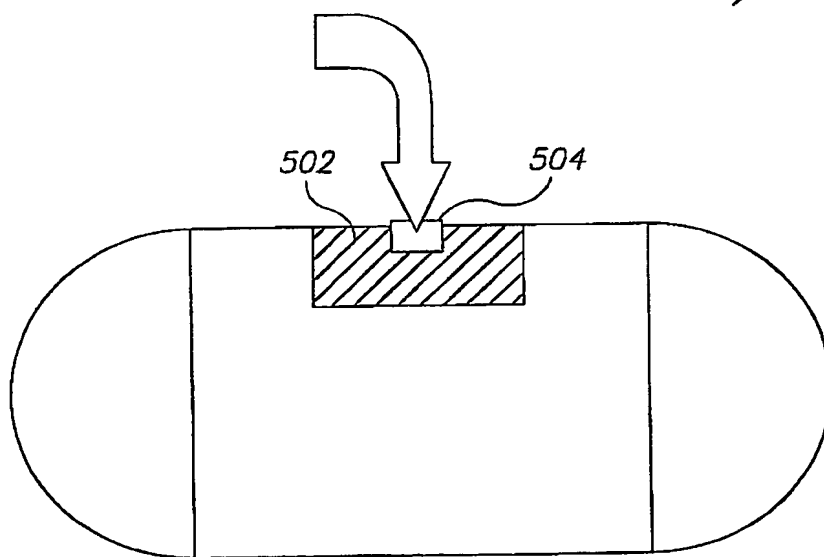


FIG. 5B

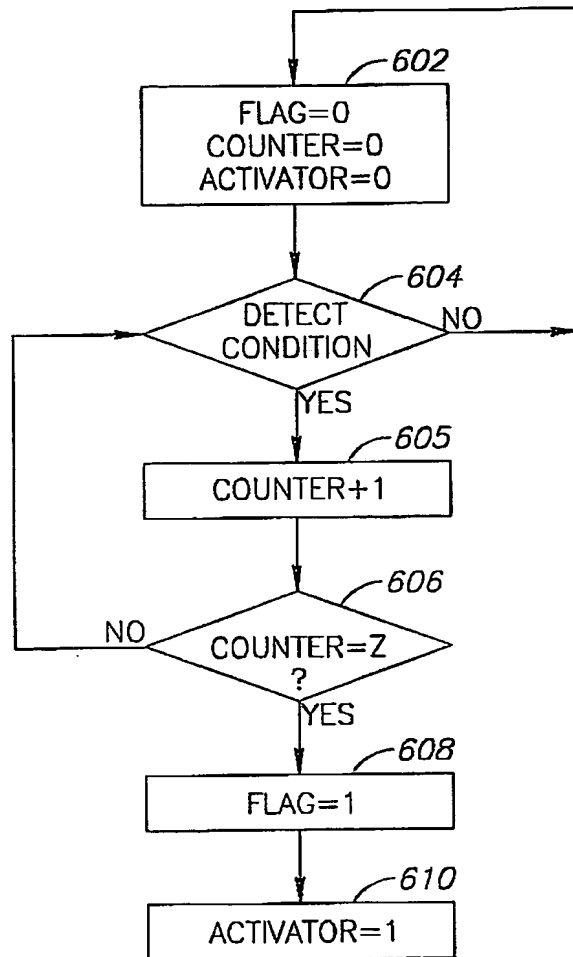


FIG. 6

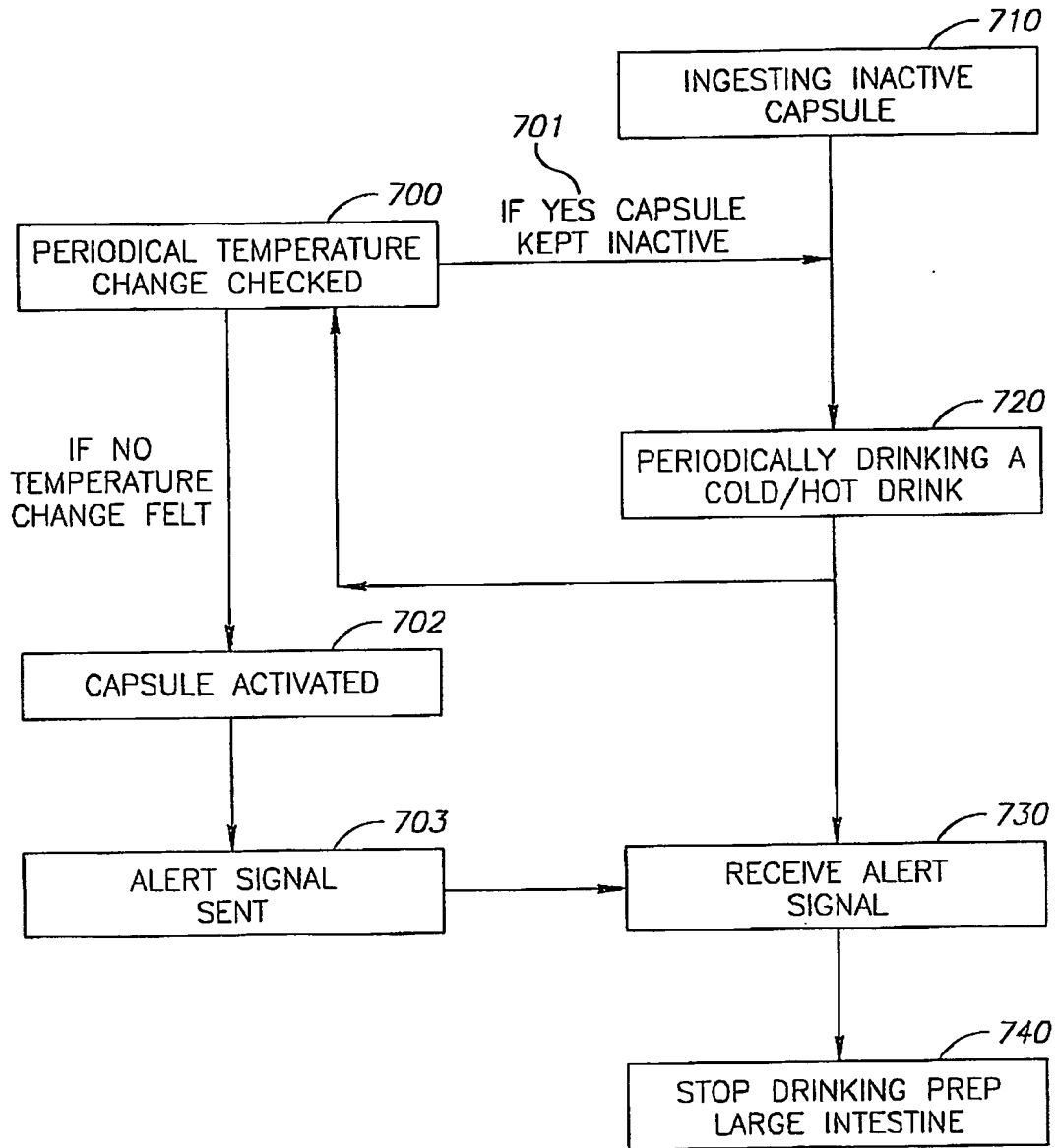


FIG.7

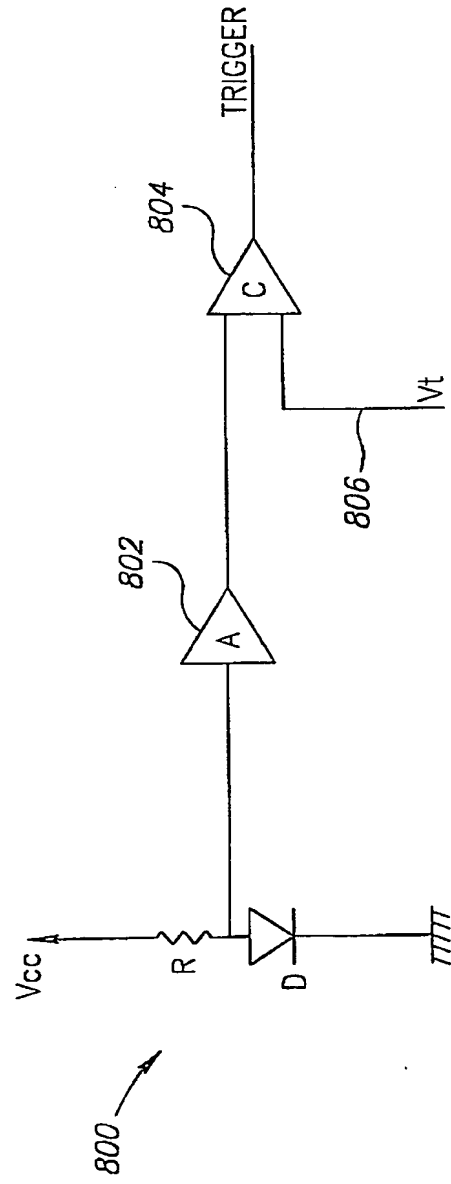


FIG.8

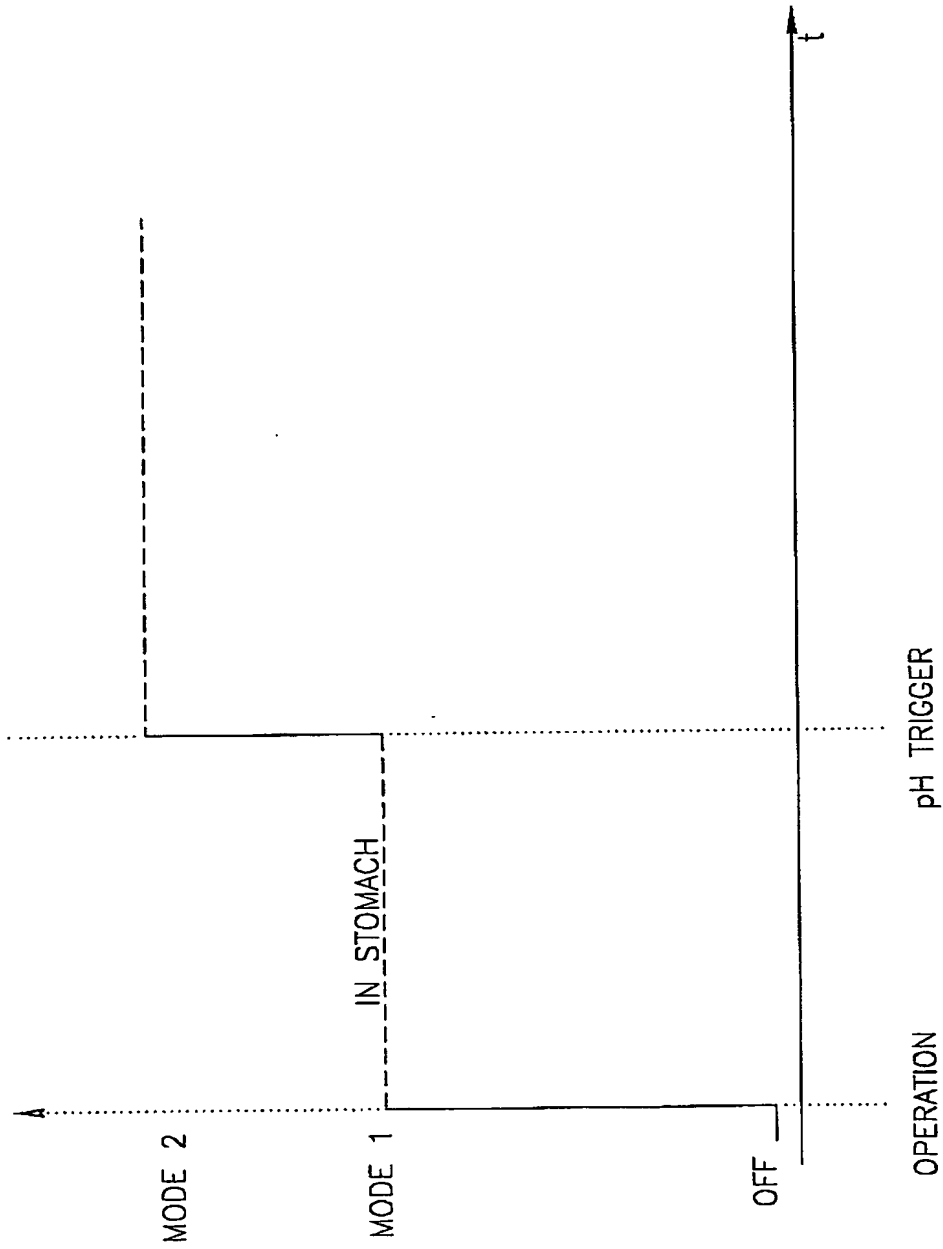


FIG.9

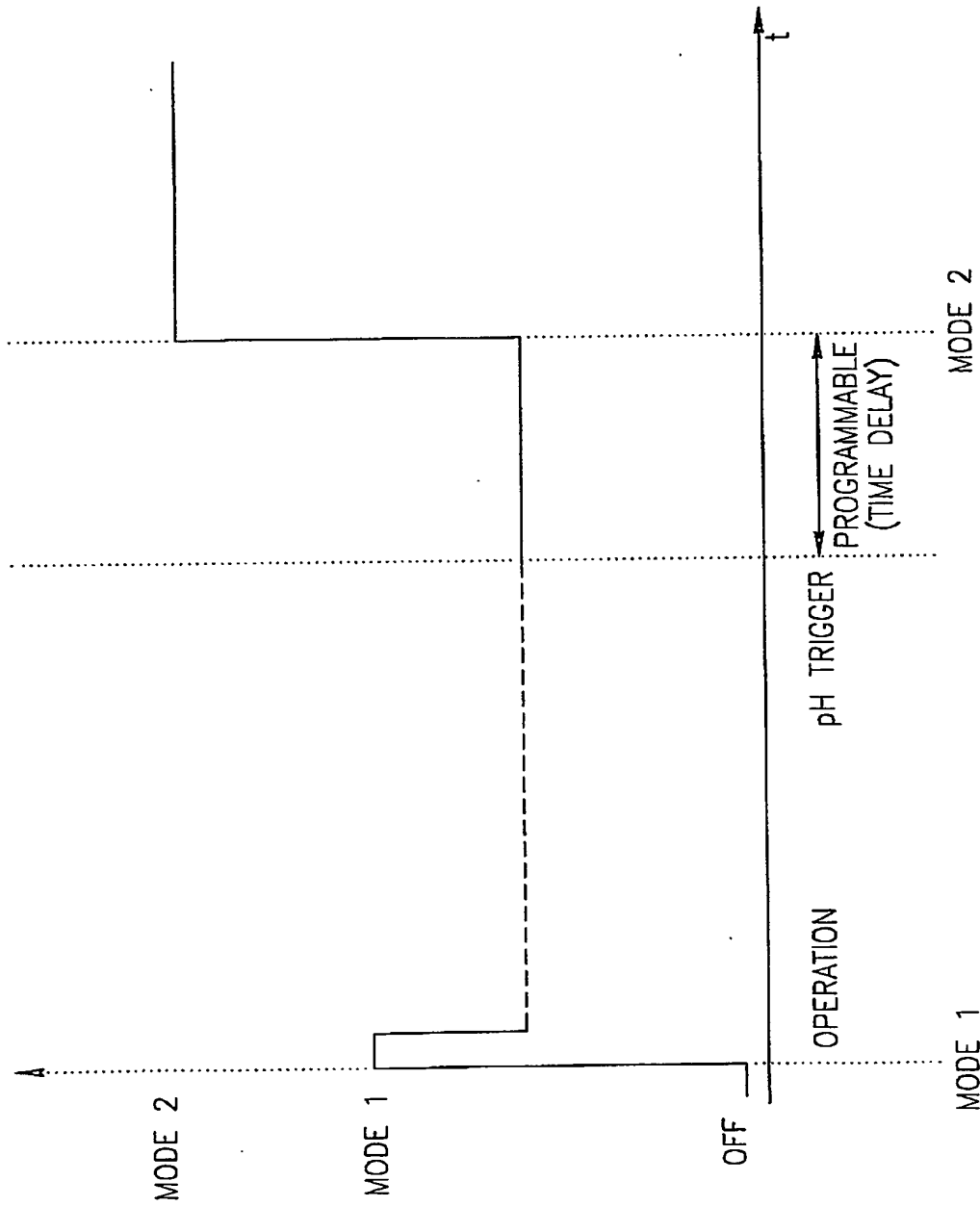


FIG.10

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

Applicant(s): GLUKHOVSKY, Arkady et al. Examiner: UNASSIGNED

Serial No.: UNASSIGNED Group Art Unit: UNASSIGNED

Filed: HEREWITH

Title: DEVICE, SYSTEM AND METHOD FOR SELECTIVE ACTIVATION OF IN VIVO SENSORS

PRELIMINARY AMENDMENT

Mail Stop Non-Fee Amendment Commissioner for Patents
P. O. Box 1450
Alexandria, VA 22313-1450

Sir:

Upon filing, and prior to Examination, kindly amend the above-identified application as follows:

Amendments to the Specification begin on page 2 of this Amendment.

Amendments to the Drawings begin on page 4 of this paper and include an attached replacement sheet.

Remarks/Arguments begin on page 5 of this paper.

An **Appendix** including amended drawing Fig. 1A is attached following page 5 of this paper.

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Page 2

AMENDMENTS TO SPECIFICATION

In the Specification:

Please add the following new paragraph after the title, before the "Field of the invention":

--CROSS-REFERENCE TO RELATED APPLICATIONS

This application is a National Phase Application of PCT International Application No. PCT/IL03/01080, International Filing Date 16 December, 2003, entitled "DEVICE, SYSTEM AND METHOD FOR SELECTIVE ACTIVATION OF IN VIVO SENSORS", in turn claiming priority of US Patent Application 60/433,586, filed 16 December, 2002, entitled "DEVICE, SYSTEM AND METHOD FOR SELECTIVE ACTIVATION OF IN VIVO SENSORS", each application incorporated herein by reference in its entirety.--

Please replace the paragraph at page 13, lines 9-29, with the following amended paragraph:

--In certain embodiments, a range of color sensitive pixels, some of which may be sensitive to the various colors that can appear on material 202 may be situated on the pixel array 210 of image sensor 46. Signals produced by each of such specific pixels 206 may vary depending on the color appearing on material 202. Controller 48 may detect and differentiate between such various signals, for example by utilizing appropriate image processing algorithms, and issue instructions to a sensor in response to each thereof. According to one embodiment a change of color may be detected in the in vivo environment that is being imaged. For example, a spot of bleeding may appear in a certain image. The change of color, that may indicate, for example, pathology in the GI tract, may be recognized by known methods. For example, controller 48, or data processor 14 may generate a probability indication of presence of colorimetric abnormalities on comparison of color content of the

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images and at least one reference value, for example, as described in PCT publication WO 02/073507, published on 19 September, 2002, that is assigned to the common assignee of the present invention. According to some embodiments, once a color change may be detected the controller 48 or data processor 14 may initiate a change in the mode of operation of device 40, of the external receiver 12, of both or of any other component or combination of components of the system. In other embodiments, a photodiode maybe used to detect changes in material 202. Such photodiode may in certain embodiments be connected to an amplifier that may be further connected to a comparator. A mode change may thereby be triggered by analog rather than digital electronics.--

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AMENDMENTS TO THE DRAWINGS

The attached sheet of drawings includes changes to Fig. 1A. This sheet, which includes Figs. 1A and 1B, replaces the original sheet including Figs. 1A and 1B. In Figure 1A, previously omitted element 44 has been added.

Attachment: Replacement Sheet

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REMARKS

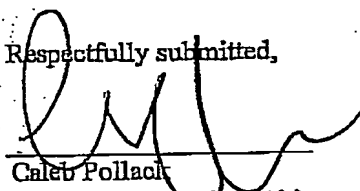
Applicants request entry of the Preliminary Amendment.

Applicants have added a cross reference to claim priority from PCT International Application No. PCT/IL03/01080. Applicants have removed an incorrect reference number (210) from the specification, as indicated above. Applicants have added previously omitted element 44 to Fig. 1A, as indicated above. No new matter has been added by the amendments to the specification or the drawings.

Should the Examiner have any question or comment as to the form, content, or entry of this Amendment, the Examiner is requested to contact the undersigned at the telephone number below.

No fees are due, however, if any fee is due, the undersigned hereby authorizes the United States Patent and Trademark Office to charge the fees to Deposit Account 05-0649.

Respectfully submitted,



Caleb Pollack
Attorney for Applicant(s)
Registration No. 37,912

Dated: April 19, 2004

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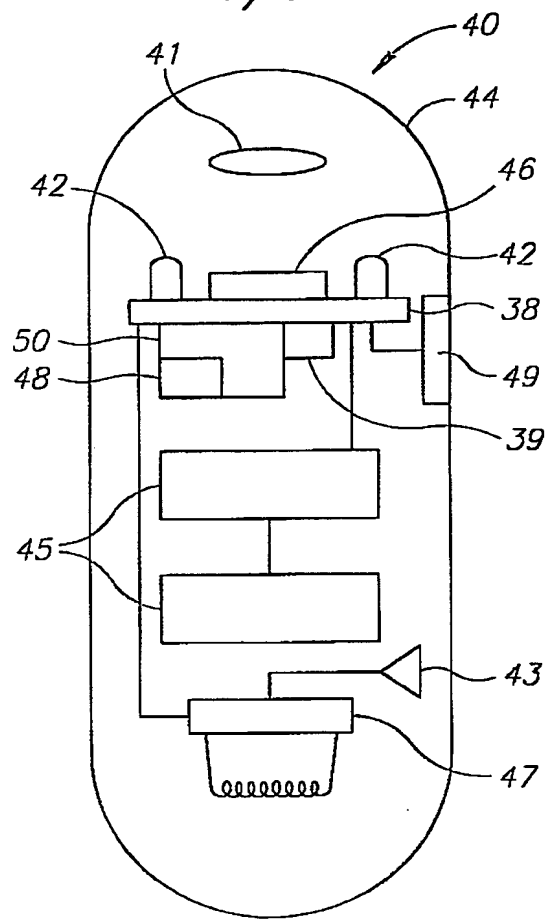


FIG. 1A

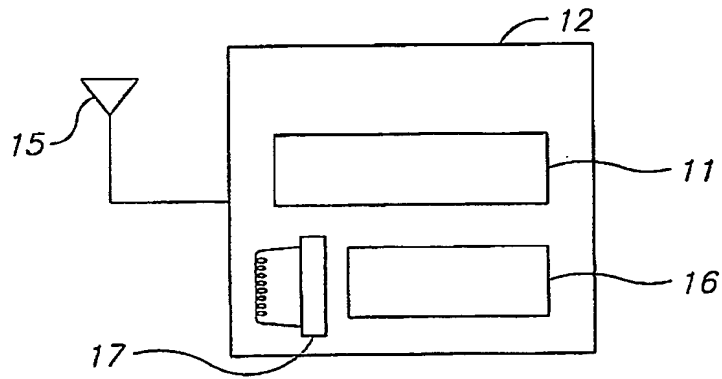


FIG. 1B

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